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Q11. N36 M

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Bansal, Geetha

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Friday, November 10, 2000 10:56 AM

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McCormick A A; Kumagai M H; Hanley K; Turpen T H; Hakim I; Grill L K; Tuse
D; Levy S; Levy R
PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF
AMERICA, (1999 Jan 19) 96 (2) 703-8

 King C A; Spellerberg M B; Zhu D; Rice J; Sahota S S; Thompsett A R; Hamblin T J; Radl J; Stevenson F K NATURE MEDICINE, (1998 Nov) 4 (11) 1281-6.

3. ***Idiotypic*** vaccination in B-cell malignancies. Bianchi A.; Massaia M. Molecular Medicine Today, (1997) 3/10 (435-441).

4. Stevenson F K; Zhu D; King C A; Ashworth L J; Kumar S; Thompsett A; Hawkins R E ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1995 Nov 27) 772 212-26.

Thanks

Geetha Bansal

1642 CM1/8A03 305-3955

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Molecular Medicine Today, (1997) 3/10 (435-441).

4. Stevenson F K; Zhu D; King C A; Ashworth L J; Kumar S; Thompsett A; Hawkins R E ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1995 Nov 27) 772 212-26.

Thanks

Geetha Bansal

1642 CM1/8A03 305-3955



NOS

Bansal, Geetha Friday, November 10, 2000 10:56 AM STIC-ILL <u>911. NJ</u> 319734

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McCormick A A; Kumagai M H; Hanley K; Turpen T H; Hakim I; Grill L K; Tuse
D; Levy S; Levy R
PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF
AMERICA, (1999 Jan 19) 96 (2) 703-8

2. King C A; Spellerberg M B; Zhu D; Rice J; Sahota S S; Thompsett A R; Hamblin T J; Radl J; Stevenson F K NATURE MEDICINE, (1998 Nov) 4 (11) 1281-6.

3. ***Idiotypic*** vaccination in B-cell malignancies. Bianchi A.; Massaia M. Molecular Medicine Today, (1997) 3/10 (435-441).

 Stevenson F K; Zhu D; King C A; Ashworth L J; Kumar S; Thompsett A; Hawkins R E ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1995 Nov 27) 772 212-26.

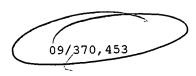
Thanks

Geetha Bansal

1642 CM1/8A03 305-3955 Agl-500 N484

1

2



FILE 'HOME' ENTERED AT 10:14:06 ON 10 NOV 2000

=> file medline biosis embase scisearch uspatfull wpids

COST IN U.S. DOLLARS

SINCE FILE TOTAL SESSION 0.75 0.75

FULL ESTIMATED COST

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FILE 'WPIDS' ENTERED AT 10:16:46 ON 10 NOV 2000 COPYRIGHT (C) 2000 DERWENT INFORMATION LTD

=> s B cell lymphoma derived immunoglobulin!

L1 0 B CELL LYMPHOMA DERIVED IMMUNOGLOBULIN!

=> s (vaccine!) and (B cell lymphoma)

L2 256 (VACCINE!) AND (B CELL LYMPHOMA)

=> s 12 and antibod?

L3 164 L2 AND ANTIBOD?

=> s 13 and multivalent vaccine

L4 4 L3 AND MULTIVALENT VACCINE

=> s 13 and (multivalent vaccine)

L5 4 L3 AND (MULTIVALENT VACCINE)

=> d 15 1-4 bib ab

L5 ANSWER 1 OF 4 USPATFULL

AN 2000:47032 USPATFULL

TI Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses

IN Rose, Timothy M., 5045 NE. 70th St., Seattle, WA, United States 98115 Bosch, Marnix L., 2601 78th Ave. NE., Bellevue, WA, United States

98004

Strand, Kurt, 22101 SE. 32 St., Issaquah, WA, United States 98027

PI US 6051375 20000418

US 1999-301390 19990428 (9) ΑI Division of Ser. No. US 1996-720229, filed on 26 Sep 1996 RLI PRAI US 1995-4297 19950926 (60) DT Utility EXNAM Primary Examiner: Mosher, Mary E.; Assistant Examiner: Salimi, Ali R. LREP Fish & Richardson, P.C. CLMN Number of Claims: 3 ECL Exemplary Claim: 1 DRWN 32 Drawing Figure(s); 33 Drawing Page(s) LN.CNT 7446 CAS INDEXING IS AVAILABLE FOR THIS PATENT. This invention relates to polynucleotides encoding Glycoprotein B from the RFHV/KSHV subfamily of gamma herpes viruses, three members of which are characterized in detail. DNA extracts were obtained from Macaque nemestrina and Macaque mulatta monkeys affected with retroperitoneal fibromatosis (RF), and human AIDS patients affected with Kaposi's sarcoma (KS). The extracts were amplified using consensus-degenerate oligonucleotide probes designed from known protein and DNA sequences of gamma herpes viruses. The nucleotide sequences of a 319 base pair fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided to characterize Glycoprotein B encoding regions of members of the RFHV/KSHV subfamily, including but not limited to RFHV1, RFHV2, and KSHV Peptides, polynucleotides, and antibodies of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B. L5ANSWER 2 OF 4 USPATFULL 2000:15318 USPATFULL ΑN ΤI Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses Rose, Timothy M., Seattle, WA, United States IN Bosch, Marnix L., Seattle, WA, United States Strand, Kurt, Issaquah, WA, United States University of Washington, Seattle, WA, United States (U.S. corporation) PΑ US 6022542 20000208 PΙ US 1996-720229 19960926 (8) ΑI US 1995-4297 19950926 (60) PRAI Utility DTPrimary Examiner: Mosher, Mary E.; Assistant Examiner: Salimi, Ali EXNAM Fish & Richardson P.C. LREP CLMN Number of Claims: 7 Exemplary Claim: 1 ECL 40 Drawing Figure(s); 33 Drawing Page(s) DRWN LN.CNT 6825 CAS INDEXING IS AVAILABLE FOR THIS PATENT. This invention relates to polynucleotides encoding Glycoprotein B from AB the RFHV/KSHV subfamily of gamma herpes viruses, three members of which are characterized in detail. DNA extracts were obtained from Macaque nemestrina and Macaque mulatta monkeys affected with retroperitoneal fibromatosis (RF), and human AIDS patients affected with Kaposi's sarcoma (KS). The extracts were amplified using consensus-degenerate oligonucleotide probes designed from known protein and DNA sequences of gamma herpes viruses. The nucleotide sequences of a 319 base pair fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside

the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided

to

characterize Glycoprotein B encoding regions of members of the $\ensuremath{\mathsf{RFHV/KSHV}}$

polynucleotides, and **antibodies** of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B.

L5 ANSWER 3 OF 4 USPATFULL

AN 1999:171946 USPATFULL

TI Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses

IN Rose, Timothy M., Seattle, WA, United States Bosch, Marnix L., Bellevue, WA, United States Strand, Kurt, Issaquah, WA, United States

PA The University of Washington, Seattle, WA, United States (U.S. corporation)

PI US 6015565 19990118

AI US 1997-804439 19970221 (8)

RLI Continuation-in-part of Ser. No. WO 1996-US15702, filed on 26 Sep 1996 And a continuation-in-part of Ser. No. US 1996-720229, filed on 26 Sep 1996

PRAI US 1995-4297 19950926 (60) US 1996-1148 19960711 (60)

DT Utility

EXNAM Primary Examiner: Eisenschenk, Frank C.; Assistant Examiner: Salimi, Ali

R.

LREP Wetherell, Jr., JohnFish & Richardson P.C.

CLMN Number of Claims: 17 ECL Exemplary Claim: 1

DRWN 33 Drawing Figure(s); 34 Drawing Page(s)

LN.CNT 7515

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

This invention relates to polynucleotides encoding Glycoprotein B from the RFHV/KSHV subfamily of gamma herpes viruses, three members of which are characterized in detail. DNA extracts were obtained from Macaque nemestrina and Macaque mulatta monkeys affected with retroperitoneal fibromatosis (RF), and human AIDS patients affected with Kaposi's sarcoma (KS). The extracts were amplified using consensus-degenerate oligonucleotide probes designed from known protein and DNA sequences of gamma herpes viruses. The nucleotide sequences of a 319 base pair fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The</pre> full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided

to

characterize Glycoprotein B encoding regions of members of the RFHV/KSHV

subfamily, including but not limited to RFHV1, RFHV2, and KSHV Peptides,

polynucleotides, and **antibodies** of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B.

```
ANSWER 4 OF 4 USPATFULL
L5
ΑN
       1999:132226 USPATFULL
       Vaccines for treatment of lymphoma and leukemia
ΤI
IN
       Denney, Jr., Dan W., Redwood City, CA, United States
       Genitope Corporation, United States (U.S. corporation)
PΑ
       US 5972334 19991026
PΙ
       US 1996-761277 19961206 (8)
ΑI
       Continuation-in-part of Ser. No. US 1996-644664, filed on 1 May 1996,
RLI
       now patented, Pat. No. US 5776746
DT
       Utility
       Primary Examiner: Hutzell, Paula K.; Assistant Examiner: Bansal, Geetha
EXNAM
       Ρ.
       Medlen & Carroll, LLP
LREP
       Number of Claims: 47
CLMN
ECL
       Exemplary Claim: 1
       26 Drawing Figure(s); 26 Drawing Page(s)
DRWN
LN.CNT 5339
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides multivalent vaccines for the
       treatment of B-cell malignancies (e.g., lymphomas and leukemias). The
       present invention also provides methods for the production of custom
     vaccines, including multivalent vaccines for the
       treatment of immune cell tumors malignancies as well as methods of
       treating immune cell tumors using custom vaccines.
=> dhis
DHIS IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).
=> d his
     (FILE 'HOME' ENTERED AT 10:14:06 ON 10 NOV 2000)
     FILE 'MEDLINE, BIOSIS, EMBASE, SCISEARCH, USPATFULL, WPIDS' ENTERED AT
     10:16:46 ON 10 NOV 2000
              O S B CELL LYMPHOMA DERIVED IMMUNOGLOBULIN!
L1
            256 S (VACCINE!) AND (B CELL LYMPHOMA)
L2
            164 S L2 AND ANTIBOD?
L3
              4 S L3 AND MULTIVALENT VACCINE
T.4
L5
              4 S L3 AND (MULTIVALENT VACCINE)
=> s 12 and (idiotyp?)
           151 L2 AND (IDIOTYP?)
=> s 13 and 16
           116 L3 AND L6
L7
=> s 17 and (recombinant V region!)
             O L7 AND (RECOMBINANT V REGION!)
L8
=> s 17 and (immunoglobulin!)
            38 L7 AND (IMMUNOGLOBULIN!)
=> dup rem 19
```

=> d 110 1-35 bib ab

```
L10 ANSWER 1 OF 35 USPATFULL
AN
      2000:141878 USPATFULL
TI
      Recombinant anti-CD4 antibodies for human therapy
IN
      Hanna, Nabil, Olivenhain, CA, United States
      Newman, Roland Anthony, San Diego, CA, United States
      Reff, Mitchell Elliot, San Diego, CA, United States
PΑ
      IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S.
      corporation)
PΙ
      US 6136310 20001024
      US 1995-523894 19950906 (8)
ΑI
      Continuation-in-part of Ser. No. US 1995-476237, filed on 7 Jun 1995,
RLI
      now patented, Pat. No. US 5756096 which is a continuation-in-part of
      Ser. No. US 1995-379072, filed on 25 Jan 1995, now patented, Pat. No.
US
      5658570 which is a continuation of Ser. No. US 1992-912292, filed on 10
      Jul 1992, now abandoned which is a continuation-in-part of Ser. No. US
      1992-856281, filed on 23 Mar 1992, now abandoned which is a
      continuation-in-part of Ser. No. US 1991-735064, filed on 25 Jul 1991,
      now abandoned
      Utility
DT
EXNAM
      Primary Examiner: Bansal, Geetha P.
LREP
      Burns, Doane, Swecker & Mathis, LLP
      Number of Claims: 16
CLMN
ECL
      Exemplary Claim: 1
DRWN
      32 Drawing Figure(s); 32 Drawing Page(s)
LN.CNT 3398
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      Chimeric antibodies specific to human CD4 antigen, DNA
      encoding, pharmaceutical compositions containing and use thereof as
      therapeutic agents are taught. These chimeric antibodies
      contain Old World monkey variable sequences and human constant domain
      sequences, preferably human gamma 1, gamma 4 or mutated forms thereof.
      These antibodies possess desirable therapeutic properties
      including low antigenicity, reduced (or absent) T cell depleting
      activity, good affinity to human CD4 and enhanced stability (in vivo
      half-life).
L10 ANSWER 2 OF 35 USPATFULL
       2000:101879 USPATFULL
ΑN
TI
      Enhancement of B cell lymphoma and tumor
       resistance using idiotype/cytokine conjugates
      Levy, Ronald, Stanford, CA, United States
IN
      Tao, Mi-Hua, Taipei, Taiwan, Province of China
      The Board of Trustees of the Leland Stanford Junior University, Palo
PΑ
      Alto, CA, United States (U.S. corporation)
ΡI
      US 6099846 20000808
      WO 9408601
                  19940428
      US 1995-416787 19950414 (8)
ΑI
      WO 1993-US9895 19931014
              19950414 PCT 371 date
              19950414 PCT 102(e) date
      Continuation-in-part of Ser. No. US 1992-961788, filed on 14 Oct 1992,
RLI
      now abandoned
      Utility
DT
      Primary Examiner: Chan, Christina Y.; Assistant Examiner: Nolan,
EXNAM
Patrick
       J.
```

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LREP
       Morrison & Foerster LLP
CLMN
       Number of Claims: 7
ECL
       Exemplary Claim: 1
DRWN
       15 Drawing Figure(s); 10 Drawing Page(s)
LN.CNT 520
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       B cell lymphoma tumor-associated antigen
AB
       or a fragment thereof containing an epitope are linked to an
       immune-enhancing cytokine, such as GM-CSF, IL-2, or IL-4 to form an
       immuno-complex. This immuno-complex elicits immune responses which are
       protective with respect to tumor proliferation. The linkers may be
       simple chemical bifunctional moieties introduced through chemical
       synthetic techniques or peptides introduce through recombinant
       methodologies. Antibodies immunoreactive with these
       immunocomplexes are also useful as passive vaccines and as
       analytical tools.
L10 ANSWER 3 OF 35 USPATFULL
       2000:54204 USPATFULL
AN
       Variable heavy and light chain regions of murine monoclonal
TI
     antibody 1F7
       Muller, Sybille, Lexington, KY, United States
IN
       Kohler, Heinz, Lexington, KY, United States
       Immpheron, Inc., Lexington, KY, United States (U.S. corporation)
PΆ
       US 6057421 20000502
PΙ
       US 1997-984277 19971203 (8)
ΑI
       Continuation-in-part of Ser. No. US 1994-351193, filed on 30 Nov 1994,
RLI
       now abandoned
DT
       Utility
      Primary Examiner: Burke, Julie
EXNAM
LREP
       Meadows, James H.
       Number of Claims: 4
CLMN
ECL
       Exemplary Claim: 1
       28 Drawing Figure(s); 23 Drawing Page(s)
DRWN
LN.CNT 2137
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The amino acid sequences of variable heavy and variable light domains
AB
of
       murine monoclonal antibody 1F7 are reported. Methods of use
       for products containing these sequences in the diagnosis and the
       treatment of HIV infection and AIDS are also described.
L10 ANSWER 4 OF 35 USPATFULL
       2000:47032 USPATFULL
AN
       Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses
TΤ
       Rose, Timothy M., 5045 NE. 70th St., Seattle, WA, United States 98115
ΤN
       Bosch, Marnix L., 2601 78th Ave. NE., Bellevue, WA, United States
98004
       Strand, Kurt, 22101 SE. 32 St., Issaquah, WA, United States 98027
       US 6051375 20000418
PΤ
       US 1999-301390 19990428 (9)
ΑI
       Division of Ser. No. US 1996-720229, filed on 26 Sep 1996 °
RLI
PRAI
       US 1995-4297
                           19950926 (60)
       Utility
DT
EXNAM Primary Examiner: Mosher, Mary E.; Assistant Examiner: Salimi, Ali R.
       Fish & Richardson, P.C.
LREP
       Number of Claims: 3
CLMN
       Exemplary Claim: 1
ECL
       32 Drawing Figure(s); 33 Drawing Page(s)
DRWN
LN.CNT 7446
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       This invention relates to polynucleotides encoding Glycoprotein B from
AB
       the RFHV/KSHV subfamily of gamma herpes viruses, three members of which
```

are characterized in detail. DNA extracts were obtained from Macaque

nemestrina and Macaque mulatta monkeys affected with retroperitoneal fibromatosis (RF), and human AIDS patients affected with Kaposi's sarcoma (KS). The extracts were amplified using consensus-degenerate oligonucleotide probes designed from known protein and DNA sequences of gamma herpes viruses. The nucleotide sequences of a 319 base pair fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided

t.o

characterize Glycoprotein B encoding regions of members of the $\ensuremath{\mathsf{RFHV/KSHV}}$

subfamily, including but not limited to RFHV1, RFHV2, and KSHV Peptides,

polynucleotides, and **antibodies** of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B.

L10 ANSWER 5 OF 35 USPATFULL

AN 2000:24287 USPATFULL

TI Receptor specific transepithelial transport of therapeutics

IN Blumberg, Richard S., Chestnut Hill, MA, United States Simister, Neil E., Wellesley, MA, United States Lencer, Wayne I., Jamaica Plain, MA, United States

PA The Brigham and Women's Hospital, Inc., Boston, MA, United States (U.S. corporation)

Brandeis University, Waltham, MA, United States (U.S. corporation)

PI US 6030613 20000229

AI US 1997-899856 19970724 (8)

Continuation-in-part of Ser. No. US 1995-578171, filed on 29 Dec 1995 which is a continuation-in-part of Ser. No. US 1995-374159, filed on 17 Jan 1995, now patented, Pat. No. US 5671273

DT Utility

EXNAM Primary Examiner: Cunningham, Thomas M.

LREP Wolf, Greenfield & Sacks, P.C.

CLMN Number of Claims: 34

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 1591

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates in general to methods and products for initiating an immune response against an antigen, and in particular relates to transepithelial delivery of antigens to provoke tolerance

and

immunity. The present invention further relates to methods and products for the transepithelial delivery of therapeutics. In particular, the invention relates to methods and compositions for the delivery of therapeutics conjugated to a FcRn binding partner to intestinal epithelium, mucosal epithelium and epithelium of the lung. The present invention further relates to the synthesis, preparation and use of the FcRn binding partner conjugates as, or in, pharmaceutical compositions for oral systemic delivery of drugs and vaccines.

L10 ANSWER 6 OF 35 USPATFULL

AN 2000:15318 USPATFULL

TI Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses

IN Rose, Timothy M., Seattle, WA, United States Bosch, Marnix L., Seattle, WA, United States Strand, Kurt, Issaquah, WA, United States

PA University of Washington, Seattle, WA, United States (U.S. corporation)

US 6022542 20000208 US 1996-720229 19960926 (8) . ΑI 19950926 (60) PRAI US 1995-4297 DTUtility Primary Examiner: Mosher, Mary E.; Assistant Examiner: Salimi, Ali EXNAM LREP Fish & Richardson P.C. Number of Claims: 7 CLMN ECL Exemplary Claim: 1 40 Drawing Figure(s); 33 Drawing Page(s) DRWN LN.CNT 6825 CAS INDEXING IS AVAILABLE FOR THIS PATENT. This invention relates to polynucleotides encoding Glycoprotein B from AB the RFHV/KSHV subfamily of gamma herpes viruses, three members of which are characterized in detail. DNA extracts were obtained from Macaque nemestrina and Macaque mulatta monkeys affected with retroperitoneal fibromatosis (RF), and human AIDS patients affected with Kaposi's sarcoma (KS). The extracts were amplified using consensus-degenerate oligonucleotide probes designed from known protein and DNA sequences of gamma herpes viruses. The nucleotide sequences of a 319 base pair fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The</pre> full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided to characterize Glycoprotein B encoding regions of members of the RFHV/KSHV subfamily, including but not limited to RFHV1, RFHV2, and KSHV Peptides, polynucleotides, and antibodies of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B. ANSWER 7 OF 35 USPATFULL L10 2000:12608 USPATFULL ΑN Methods for determining the presence of carcinoma using the antigen ΤI binding region of monoclonal antibody BR96 Hellstrom, Ingegerd, Seattle, WA, United States IN Hellstrom, Karl Erik, Seattle, WA, United States Bruce, Kim Folger, Seattle, WA, United States Schreiber, George J., Seattle, WA, United States Bristol-Myers Squibb Company, Princeton, NJ, United States (U.S. PA corporation) US 6020145 20000201 PIUS 1994-333840 19941103 (8) ΑI Division of Ser. No. US 1993-77253, filed on 14 Jun 1993 which is a RLI continuation-in-part of Ser. No. US 1993-57444, filed on 5 May 1993, now patented, Pat. No. US 5491088 which is a continuation of Ser. No. US 1990-544246, filed on 26 Jun 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-374947, filed on 30 Jun 1989, now abandoned DT Utility Primary Examiner: Feisee, Lila; Assistant Examiner: Bansal, Geetha P. EXNAM Merchant, Gould, Smith, Edell, Welter & Schmidt LREP CLMN Number of Claims: 4 ECLExemplary Claim: 1,3 76 Drawing Figure(s); 74 Drawing Page(s) LN.CNT 5875 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention relates to novel antibodies,

```
antibody fragments and antibody conjugates and
       single-chain immunotoxins reactive with human carcinoma cells. More
      particularly, the antibodies, conjugates and single-chain
       immunotoxins of the invention include: a murine monoclonal
     antibody, BR96; a human/murine chimeric antibody,
       ChiBR96; a F(ab').sub.2 fragment of BR96; ChiBR96-PE, ChiBR96-LysPE40,
       ChiBR96 F(ab').sub.2 -LysPE40 and ChiBR96 Fab'-LysPE40 conjugates and
       recombinant BR96 sFv-PE40 immunotoxin. These molecules are reactive
with
      a cell membrane antigen on the surface of human carcinomas. The BR96
     antibody and its functional equivalents, displays a high degree
      of selectivity for carcinoma cells and possess the ability to mediate
     antibody-dependent cellular cytotoxicity and
       complement-dependent cytotoxicity activity. In addition, the
     antibodies of the invention internalize within the carcinoma
      cells to which they bind and are therefore particularly useful for
      therapeutic applications, for example, as the antibody
      component of antibody-drug or antibody-toxin
      conjugates. The antibodies also have a unique feature in that
       they are cytotoxic when used in the unmodified form, at specified
      concentrations.
L10 ANSWER 8 OF 35 USPATFULL
       1999:171946 USPATFULL
ΑN
TI
       Glycoprotein B of the RFHV/KSHV subfamily of herpes viruses
      Rose, Timothy M., Seattle, WA, United States
IN
      Bosch, Marnix L., Bellevue, WA, United States
      Strand, Kurt, Issaquah, WA, United States
      The University of Washington, Seattle, WA, United States (U.S.
PΑ
      corporation)
      US 6015565 19990118
PΙ
      US 1997-804439 19970221 (8)
ΑI
      Continuation-in-part of Ser. No. WO 1996-US15702, filed on 26 Sep 1996
RLI
      And a continuation-in-part of Ser. No. US 1996-720229, filed on 26 Sep
      1996
      US 1995-4297
                           19950926 (60)
PRAI
      US 1996-1148
                           19960711 (60)
DT
      Utility
      Primary Examiner: Eisenschenk, Frank C.; Assistant Examiner: Salimi,
EXNAM
Ali
LREP
      Wetherell, Jr., JohnFish & Richardson P.C.
CLMN
      Number of Claims: 17
ECL
      Exemplary Claim: 1
       33 Drawing Figure(s); 34 Drawing Page(s)
DRWN
LN.CNT 7515
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       This invention relates to polynucleotides encoding Glycoprotein B from
       the RFHV/KSHV subfamily of gamma herpes viruses, three members of which
       are characterized in detail. DNA extracts were obtained from Macaque
       nemestrina and Macaque mulatta monkeys affected with retroperitoneal
       fibromatosis (RF), and human AIDS patients affected with Kaposi's
       sarcoma (KS). The extracts were amplified using consensus-degenerate
       oligonucleotide probes designed from known protein and DNA sequences of
       gamma herpes viruses. The nucleotide sequences of a 319 base pair
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fragment are about 76% identical between RFHV1 and KSHV, and about 60-63% identical with the closest related gamma herpes viruses outside

the RFHV/KSHV subfamily. Protein sequences encoded within these fragments are are about 91% identical between RFHV1 and KSHV, and <.about.65% identical to that of other gamma herpes viruses. The full-length KSHV Glycoprotein B sequence comprises a transmembrane domain near the N-terminus, and a plurality of potentially antigenic sites in the extracellular domain. Materials and methods are provided

characterize Glycoprotein B encoding regions of members of the RFHV/KSHV subfamily, including but not limited to RFHV1, RFHV2, and KSHV polynucleotides, and antibodies of this invention can be used for diagnosing infection, and for eliciting an immune response against Glycoprotein B. ANSWER 9 OF 35 USPATFULL L10 1999:141303 USPATFULL ANAntibodies reactive with human carcinomas TIHellstrom, Ingegerd, Seattle, WA, United States IN Hellstrom, Karl Erik, Seattle, WA, United States Bruce, Kim Folger, Seattle, WA, United States Schreiber, George J., Redmond, WA, United States Siegall, Clay, Edmonds, WA, United States McAndrew, Stephen, Newtown, PA, United States Bristol-Myers Squibb Company, Princeton, NJ, United States (U.S. PΑ corporation) US 5980896 19991109 PΙ ΑI US 1993-77253 19930614 (8) Continuation-in-part of Ser. No. US 1993-57444, filed on 5 May 1993, RLI now patented, Pat. No. US 5491088 And Ser. No. US 1992-892501, filed on 1 Jun 1992, now abandoned which is a continuation-in-part of Ser. No. US 1990-544246, filed on 26 Jun 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-374947, filed on 30 Jun 1989, now abandoned , said Ser. No. US 1993-57444, filed on 5 May 1993, now patented, Pat. No. US 5491088 which is a continuation of Ser. No. US 544246 Utility DT Primary Examiner: Feisee, Lila; Assistant Examiner: Bansal, Geetha P. EXNAM Merchant, Gould, Smith, Edell, Welter & Schmidt LREP Number of Claims: 35 CLMN Exemplary Claim: 1,16,34 ECL 76 Drawing Figure(s); 74 Drawing Page(s) DRWN LN.CNT 5987 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention relates to novel antibodies, AΒ antibody fragments and antibody conjugates and single-chain immunotoxins reactive with human carcinoma cells. More particularly, the antibodies, conjugates and single-chain immunotoxins of the invention include: a murine monoclonal antibody, BR96; a human/murine chimeric antibody, ChiBR96; a F(ab').sub.2 fragment of BR96; ChiBR96-PE, ChiBR96-LysPE40, ChiBR96 F(ab').sub.2 -LysPE40 and ChiBR96 Fab'-LysPE40 conjugates and recombinant BR96 sFv-PE40 immunotoxin. These molecules are reactive with a cell membrane antigen on the surface of human carcinomas. The BR96 antibody and its functional equivalents, displays a high degree of selectivity for carcinoma cells and possess the ability to mediate antibody-dependent cellular cytotoxicity and. complement-dependent cytotoxicity activity. In addition, the antibodies of the invention internalize within the carcinoma cells to which they bind and are therefore particularly useful for therapeutic applications, for example, as the antibody component of antibody-drug or antibody-toxin conjugates. The antibodies also have a unique feature in that they are cytotoxic when used in the unmodified form, at specified concentrations. L10 ANSWER 10 OF 35 USPATFULL

AN 1999:137463 USPATFULL

TI Murine anti-idiotype antibody 3H1

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Chatterjee, Sunil K., Lexington, KY, United States
       Foon, Kenneth A., Lexington, KY, United States
       The Board of Trustees of the University of Kentucky, Lexington, KY,
PA
       United States (U.S. corporation)
РΤ
       US 5977315 19991102
       US 1995-579940 19951228 (8)
ΑI
       Continuation-in-part of Ser. No. US 1994-365484, filed on 28 Dec 1994,
RLI
       now abandoned
DT
       Utility
       Primary Examiner: Reeves, Julie
EXNAM
       Morrison & Foerster LLP
       Number of Claims: 29
CLMN
ECL
       Exemplary Claim: 1
       52 Drawing Figure(s); 43 Drawing Page(s)
DRWN
LN.CNT 2698
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides a monoclonal anti-idiotype
AB
     antibody 3H1 that escapes immune tolerance and elicits a
       specific immune response to CEA in mice, rabbits, monkeys, and patients
       with advanced CEA-associated disease. This invention also provides
       compositions which can be used in the detection or treatment of
       CEA-associated tumors mimics a specific epitope on carcinoembryonic
       antigen and a hybridoma that produces 3H1.
L10 ANSWER 11 OF 35 USPATFULL
       1999:132226 USPATFULL
AN
       Vaccines for treatment of lymphoma and leukemia
ΤI
       Denney, Jr., Dan W., Redwood City, CA, United States
ΙN
       Genitope Corporation, United States (U.S. corporation)
PΑ
       US 5972334 19991026
PΤ
ΑI
       US 1996-761277 19961206 (8)
       Continuation-in-part of Ser. No. US 1996-644664, filed on 1 May 1996,
RLI
       now patented, Pat. No. US 5776746
DT
       Utility
       Primary Examiner: Hutzell, Paula K.; Assistant Examiner: Bansal, Geetha
EXNAM
       Medlen & Carroll, LLP
LREP
CLMN
       Number of Claims: 47
       Exemplary Claim: 1
ECL
       26 Drawing Figure(s); 26 Drawing Page(s)
LN.CNT 5339
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides multivalent vaccines for the
       treatment of B-cell malignancies (e.g., lymphomas and leukemias). The
       present invention also provides methods for the production of custom
     vaccines, including multivalent vaccines for the
       treatment of immune cell tumors malignancies as well as methods of
       treating immune cell tumors using custom vaccines.
    ANSWER 12 OF 35 USPATFULL
       1999:18719 USPATFULL
ΑN
       Antibody conjugates reactive with human carcinomas
ΤI
       Hellstrom, Ingegerg, Seattle, WA, United States
TN
       Hellstrom, Karl Erik, Seattle, WA, United States
       Bruce, Kim Folger, Seattle, WA, United States
       Schreiber, George J., Seattle, WA, United States
       Bristol-Myers Squibb Company, New York, NY, United States (U.S.
PA
       corporation)
       US 5869045
                  19990209
PΤ
       US 1995-459354 19950602 (8)
AΙ
       Division of Ser. No. US 1993-77253, filed on 14 Jun 1993 which is a
RLI
       continuation-in-part of Ser. No. US 1993-57444, filed on 5 May 1993,
now
                                                                        Page 11
```

Chatterjee, Malaya, Lexington, KY, United States

Kohler, Heinz, Lexington, KY, United States

IN

patented, Pat. No. US 5491088 which is a continuation of Ser. No. US 1990-544246, filed on 26 Jun 1990, now abandoned which is a continuation-in-part of Ser. No. US 1989-374947, filed on 30 Jun 1989, now abandoned Utility Primary Examiner: Feisee, Lila; Assistant Examiner: Ungar, Susan EXNAM Merchant, Gould, Smith, Welter and Schmidt LREP Number of Claims: 7 CLMN Exemplary Claim: 1 ECL 75 Drawing Figure(s); 74 Drawing Page(s) DRWN LN.CNT 5935 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention relates to novel antibodies, AΒ antibody fragments and antibody conjugates and single-chain immunotoxins reactive with human carcinoma cells. More particularly, the antibodies, conjugates and single-chain immunotoxins of the invention include: a murine monoclonal antibody, BR96; a human/murine chimeric antibody, ChiBR96; a F(ab').sub.2 fragment of BR96; ChiBR96-PE, ChiBR96-LysPE40, ChiBR96 F(ab').sub.2 -LysPE40 and ChiBR96 Fab'-LysPE40 conjugates and recombinant BR96 sFv-PE40 immunotoxin. These molecules are reactive with a cell membrane antigen on the surface of human carcinomas. The BR96 antibody and its functional equivalents, displays a high degree of selectivity for carcinoma cells and possess the ability to mediate antibody-dependent cellular cytotoxicity and complement-dependent cytotoxicity activity. In addition, the antibodies of the invention internalize within the carcinoma cells to which they bind and are therefore particularly useful for therapeutic applications, for example, as the antibody component of antibody-drug or antibody-toxin conjugates. The antibodies also have a unique feature in that they are cytotoxic when used in the unmodified form, at specified concentrations. L10 ANSWER 13 OF 35 USPATFULL 1999:7146 USPATFULL ΑN Method and composition for transfer of active tumor-specific TΙ immunization from an immunized allogeneic bone marrow donor Kwak, Larry W., Frederick, MD, United States ΙN Longo, Dan L., Kensington, MD, United States The United States of America as represented by the Deptartment of PΑ Health and Human Services, Washington, DC, United States (U.S. government) US 5861158 19990119 PΙ US 1993-153464 19931117 (8) ΑI Utility Primary Examiner: Minnifield, Nita EXNAM Needle & Rosenberg, P.C. LREP Number of Claims: 17 CLMN ECL Exemplary Claim: 1 No Drawings DRWN LN.CNT 732 CAS INDEXING IS AVAILABLE FOR THIS PATENT. This invention provides a method of improving a transplantation of hematopoietic cells from a donor to a recipient to treat a hematopoietic cell tumor in the recipient comprising immunizing the donor's hematopoietic cells with an antigen specific for the recipient's hematopoietic cell tumor, and transplanting the donor's immunized hematopoietic cells to the recipient. Also provided is a composition comprising purified hematopoietic cells primed to produce an immunological response to foreign tumor specific antigen. Also provided is a method of treating a tumor by the transplantation of hematopoietic cells from a donor to a recipient to treat the tumor in the recipient comprising immunizing the donor's hematopoietic cells with an antigen specific for the recipient's tumor, and transplanting the donor's immunized hematopoietic cells to the recipient.

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ANSWER 14 OF 35 MEDLINE
L10
                                                         DUPLICATE 1
     1999110954
                    MEDLINE
ΑN
     99110954
DN
     Rapid production of specific vaccines for lymphoma by expression
TI
     of the tumor-derived single-chain Fv epitopes in tobacco plants.
     McCormick A A; Kumagai M H; Hanley K; Turpen T H; Hakim I; Grill L K;
ΑU
Tuse
     D; Levy S; Levy R
     Biosource Technologies, Inc., 3333 Vacavalley Parkway, Suite 1000,
CS
     Vacaville, CA 95688, USA.
     CA33399 (NCI)
NC
     AI37219 (NIAID)
     PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF
SO
     AMERICA, (1999 Jan 19) 96 (2) 703-8.
     Journal code: PV3. ISSN: 0027-8424.
ÇΥ
     United States
DT
     Journal; Article; (JOURNAL ARTICLE)
LA
     English
FS
     Priority Journals; Cancer Journals
ΕM
     199905
EW
     19990503
     Rapid production of protein-based tumor-specific vaccines for
AB ·
     the treatment of malignancies is possible with the plant-based transient
     expression system described here. We created a modified tobamoviral
vector
     that encodes the idiotype-specific single-chain Fv fragment
     (scFv) of the immunoglobulin from the 38C13 mouse B cell
     lymphoma. Infected Nicotiana benthamiana plants contain high
     levels of secreted scFv protein in the extracellular compartment. This
     material reacts with an anti-idiotype antibody by
     Western blotting, ELISA, and affinity chromatography, suggesting that the
     plant-produced 38C13 scFv protein is properly folded in solution. Mice
     vaccinated with the affinity-purified 38Cl3 scFv generate >10
     micrograms/ml anti-idiotype immunoglobulins. These
     mice were protected from challenge by a lethal dose of the syngeneic
38C13
     tumor, similar to mice immunized with the native 38C13 IgM-keyhole limpet
     hemocyanin conjugate vaccine. This rapid production system for generating
     tumor-specific protein vaccines may provide a viable strategy
     for the treatment of non-Hodgkin's lymphoma.
    ANSWER 15 OF 35 USPATFULL
L10
ΑN
       1998:147025 USPATFULL
       Vaccine comprising anti-idiotypic antibody to
TI
       chlamydia GLXA and process
IN
       MacDonald, Alex Bruce, Amherst, MA, United States
       An, Ling-Ling, La Jolla, CA, United States
       Sutton-Stuart, Elizabeth, Amherst, MA, United States
       Whittum-Hudson, Judith A., Elkton, MD, United States
       Johns Hopkins University, United States (U.S. corporation)
PΑ
       University of Massachusetts, United States (U.S. corporation)
       US 5840297 19981124
ΡI
       US 1993-34572 19930319 (8)
ΑI
\mathsf{D}\mathbf{T}
       Utility
       Primary Examiner: Loring, Susan A.
EXNAM
       Cook, Paul J.
LREP
       Number of Claims: 17
CLMN
ECL
       Exemplary Claim: 5
       17 Drawing Figure(s); 9 Drawing Page(s)
DRWN
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LN.CNT 2015
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      A genus specific chlamydia vaccine is provided which comprises an anti-
    idiotype antibody capable of producing in an animal an
      anti-anti-idiotypic antibody which recognizes a
      glycoplipid exoantigen (GLXA) of chlamydia. The vaccine is produced by
      producing an idiotypic antibody to GLXA which, in
      turn, is utilized t produce the anti-idiotypic
    antibody comprising the vaccine.
L10 ANSWER 16 OF 35 USPATFULL
AN
      1998:64956 USPATFULL
      Immunogenic cancer proteins and peptides and methods of use
TI
      Calenoff, Emanuel, Chicago, IL, United States
IN
      Northwestern University, Evanston, IL, United States (U.S. corporation)
PΑ
      US 5763164 19980609
PΙ
      US 1994-191338 19940203 (8)
ΑI
      Continuation-in-part of Ser. No. US 1993-49698, filed on 16 Apr 1993,
RLI
      now abandoned
DΨ
      Utility
      Primary Examiner: Jones, W. Gary; Assistant Examiner: Rees, Dianne
EXNAM
      Brinks Hofer Gilson & Lione
      Number of Claims: 11
CLMN
      Exemplary Claim: 1
\mathsf{ECL}
       13 Drawing Figure(s); 13 Drawing Page(s)
DRWN
LN.CNT 2928
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention relates to tumor specific antigens and functional
       proteins of a tumor cell preparable by identifying protein presents in
       the tumor cell that are selectively immunogenic for tumor patients. The
      present invention still further provides a process of making a peptide
       library of tumor specific humoral antigens, a process of increasing the
       immunogenic specificity of a tumor-associated antigen, an assay kit for
       detecting the presence of an antibody immunoreactive with a
       tumor-specific antigen, and a process of making T cells sensitized to a
       tumor-specific antigen.
L10 ANSWER 17 OF 35 USPATFULL
       1998:57523 USPATFULL
AN
       Recombinant antibodies for human therapy
ΤI
       Newman, Roland A., San Diego, CA, United States
IN
       Hanna, Nabil, Olivenhain, CA, United States
       Raab, Ronald W., San Diego, CA, United States
       IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S.
PΑ
       corporation)
       US 5756096 19980526
PΙ
       US 1995-476237 19950607 (8)
ΑI
       Continuation-in-part of Ser. No. US 1995-379072, filed on 25 Jan 1995,
RLI
       now patented, Pat. No. US 5658570 which is a continuation of Ser. No.
US
       1992-912292, filed on 10 Jul 1992, now abandoned which is a
       continuation-in-part of Ser. No. US 1992-856281, filed on 23 Mar 1992,
       now abandoned which is a continuation-in-part of Ser. No. US
       1991-735064, filed on 25 Jul 1991, now abandoned
DT
       Utility
       Primary Examiner: Feisee, Lila; Assistant Examiner: Bansal, Geetha P.
EXNAM
       Burns, Doane, Swecker & Mathis, L.L.P.
LREP
       Number of Claims: 6
CLMN
       Exemplary Claim: 1,4
ECL
       26 Drawing Figure(s); 26 Drawing Page(s)
LN.CNT 1919
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Chimeric antibodies including an Old World monkey portion and
```

a human portion, nucleic acid encoding such antibodies, Old

World monkey monoclonal antibodies, and methods for their production and use. ANSWER 18 OF 35 USPATFULL 1998:51191 USPATFULL Recombinant antibodies for human therapy Newman, Roland A., San Diego, CA, United States Hanna, Nabil, Olivenhain, CA, United States Raab, Ronald W., San Diego, CA, United States IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S. corporation) US 5750105 19980512 US 1995-476349 19950607 (8) Division of Ser. No. US 1995-379072, filed on 5 Dec 1995 which is a continuation of Ser. No. US 1992-912292, filed on 10 Jul 1992, now abandoned which is a continuation-in-part of Ser. No. US 1992-856281, filed on 23 Mar 1992, now abandoned which is a continuation-in-part of Ser. No. US 1991-735064, filed on 25 Jul 1991, now abandoned Utility Primary Examiner: Feisee, Lila; Assistant Examiner: Bansal, Geetha P. EXNAM Burns, Doane, Swecker & Mathis LLP Number of Claims: 10 Exemplary Claim: 1 26 Drawing Figure(s); 26 Drawing Page(s) LN.CNT 2110 CAS INDEXING IS AVAILABLE FOR THIS PATENT. Chimeric antibodies including an Old World monkey portion and a human portion, nucleic acid encoding such antibodies, Old World monkey monoclonal antibodies, and methods for their production and use.

L10 ANSWER 19 OF 35 USPATFULL

1998:14646 USPATFULL AN

Method for diagnosing a patient for chlamydia TI

MacDonald, Alex Bruce, Amherst, MA, United States ΙN Stuart, Elizabeth S., Amherst, MA, United States

An, Ling Ling, La Jolla, CA, United States Whipkey, Myron D., Portland, ME, United States

Animal House, Inc., Portland, ME, United States (U.S. corporation) PΑ

ΡI US 5716793 19980210

US 1995-406113 19950317 (8) ΑI

Continuation-in-part of Ser. No. US 1993-34572, filed on 19 Mar 1993 RLI

Utility

L10ΑN

TΙ

IN

PΑ

ΡI

ΑI

DT

LREP

CLMN

ECL

AΒ

DRWN

RLI

Primary Examiner: Spiegel, Carol A. EXNAM

Cook, Paul J. LREP

Number of Claims: 10 CLMN

ECL Exemplary Claim: 1

17 Drawing Figure(s); 9 Drawing Page(s)

LN.CNT 1933

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method of detecting chlamydia in a extracellular sample is provided AB which comprises contacting the sample with an idiotypic

antibody to GLXA to form an immunocomplex and detecting the immunocomplex.

ANSWER 20 OF 35 USPATFULL L10

1998:6785 USPATFULL ΑN

Induction of cytotoxic T-lymphocyte responses ΤI

Raychaudhuri, Syamal, San Diego, CA, United States IN Rastetter, William H., Rancho Santa Fe, CA, United States

IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S. PA corporation)

US 5709860 19980120 ΡI

US 1994-351001 19941207 (8) ΑI

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Continuation-in-part of Ser. No. US 1992-919787, filed on 24 Jul 1992 which is a continuation-in-part of Ser. No. US 1991-735069, filed on 25 Jul 1991, now abandoned
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DT Utility

EXNAM Primary Examiner: Woodward, Michael P.; Assistant Examiner: Zeman, Mary

LREP Burns, Doane, Swecker & Mathis, LLP

CLMN Number of Claims: 23

ECL Exemplary Claim: 1

DRWN 19 Drawing Figure(s); 14 Drawing Page(s)

LN.CNT 1242

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Methods and compositions useful for inducing a cytotoxic T lymphocyte response (CTL) in a human or domesticated or agriculturally important animal. The method includes the steps of providing the antigen to which the CTL response is desired and providing an antigen formulation which comprises, consists, or consists essentially of two or more of a stabilizing detergent, a micelle-forming agent, and an oil. This

antigen

formulation is preferably lacking in an immunostimulating peptide component, or has sufficiently low levels of such a component that the desired CTL response is not diminished. This formulation is provided as a stable oil-in-water emulsion.

- L10 ANSWER 21 OF 35 MEDLINE
- AN 1999025406 MEDLINE
- DN 99025406
- DNA vaccines with single-chain Fv fused to fragment C of tetanus toxin induce protective immunity against lymphoma and myeloma [see comments].
- CM Comment in: Nat Med 1998 Nov; 4(11):1239-40
- AU King C A; Spellerberg M B; Zhu D; Rice J; Sahota S S; Thompsett A R; Hamblin T J; Radl J; Stevenson F K
- CS Tenovus Laboratory, Southampton University Hospitals Trust, England.
- SO NATURE MEDICINE, (1998 Nov) 4 (11) 1281-6. Journal code: CG5. ISSN: 1078-8956.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 199902
- EW 19990204
- Vaccination with idiotypic protein protects against Bcell lymphoma, mainly through anti-idiotypic
 antibody. For use in patients, DNA vaccines containing
 single-chain Fv derived from tumor provide a convenient alternative
 vaccine delivery system. However, single-chain Fv sequence alone induces
 low anti-idiotypic response and poor protection against
 lymphoma. Fusion of the gene encoding fragment C of tetanus toxin to
 single-chain Fv substantially promotes the anti-idiotypic
 response and induces strong protection against B-cell
 lymphoma. The same fusion design also induces protective immunity
 against a surface Ig-negative myeloma. These findings indicate that

fusion

to a pathogen sequence allows a tumor antigen to engage diverse immune mechanisms that suppress growth. This fusion design has the added advantage of overcoming potential tolerance to tumor that may exist in patients.

- L10 ANSWER 22 OF 35 USPATFULL
- AN 97:114941 USPATFULL
- TI Induction of cytotoxic T-lymphocyte responses
- IN Raychaudhuri, Syamal, San Diego, CA, United States Rastetter, William H., Rancho Santa Fe, CA, United States

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Black, Amelia, Cardiff, CA, United States
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PA IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S. corporation)

PI US 5695770 19971209

AI US 1995-472311 19950607 (8)

RLI Continuation of Ser. No. US 1994-351001, filed on 7 Dec 1994 which is a continuation-in-part of Ser. No. US 1992-919787, filed on 24 Jul 1992, now patented, Pat. No. US 5585103, issued on 17 Dec 1996 which is a continuation-in-part of Ser. No. US 1991-735069, filed on 25 Jul 1991, now abandoned

DT Utility

EXNAM Primary Examiner: Woodward, Michael P.; Assistant Examiner: Zeman, Mary K.

LREP Burns, Doane, Swecker & Mathis, LLP

CLMN Number of Claims: 9

ECL Exemplary Claim: 1

DRWN 19 Drawing Figure(s); 14 Drawing Page(s)

LN.CNT 1134

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Methods and compositions useful for inducing a cytotoxic T lymphocyte response (CTL) in a human or domesticated or agriculturally important animal. The method includes the steps of providing the antigen to which the CTL response is desired and providing an antigen formulation which comprises, consists, or consists essentially of two or more of a stabilizing detergent, a micelle-forming agent, and an oil. This

antigen

formulation is preferably lacking in an immunostimulating peptide component, or has sufficiently low levels of such a component that the desired CTL response is not diminished. This formulation is provided as a stable oil-in-water emulsion.

L10 ANSWER 23 OF 35 USPATFULL

AN 97:112606 USPATFULL

TI Recombinant antibodies for human therapy

IN Newman, Roland A., San Diego, CA, United States Hanna, Nabil, Olivenhain, CA, United States Raab, Ronald W., San Diego, CA, United States

PA Idec Pharmaceuticals Corporation, San Diego, CA, United States (U.S. corporation)

PI US 5693780 19971202

AI US 1995-481869 19950607 (8)

Division of Ser. No. US 1995-379072, filed on 25 Jan 1995 which is a continuation of Ser. No. US 1992-912292, filed on 10 Jul 1992, now abandoned which is a continuation-in-part of Ser. No. US 1992-856281, filed on 23 Mar 1992, now abandoned which is a continuation-in-part of Ser. No. US 1991-735064, filed on 25 Jul 1991, now abandoned

DT Utility

EXNAM Primary Examiner: Scheiner, Toni R.; Assistant Examiner: Bansal, Geetha P.

LREP Burns, Doane, Swecker & Mathis, LLP

CLMN Number of Claims: 16

ECL Exemplary Claim: 1

DRWN 26 Drawing Figure(s); 26 Drawing Page(s)

LN.CNT 1755

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Chimeric antibodies including an Old World monkey portion and a human portion, nucleic acid encoding such antibodies, Old World monkey monoclonal antibodies, and methods for their production and use.

L10 ANSWER 24 OF 35 USPATFULL

AN 97:109699 USPATFULL

TI Immunoglobulin superantigen binding to gp 120 from HIV

IN ' Braun, Jonathan, Sherman Oaks, CA, United States

```
Goodglick, Lee A., Los Angeles, CA, United States
       The Regents of the University of California, Oakland, CA, United States
PΑ
      (U.S. corporation)
       US 5691135 19971125
PΙ
       US 1994-306116 19940914 (8)
ΑI
       Continuation-in-part of Ser. No. US 1994-259669, filed on 14 Jun 1994,
RLI
       now abandoned which is a continuation of Ser. No. US 1993-9705, filed
on
       26 Jan 1993, now abandoned
DT
       Utility
       Primary Examiner: Nucker, Christine M.; Assistant Examiner: Stucker,
EXNAM
       Jeffrey
       Knobbe Martens Olson & Bear, LLP
LREP
       Number of Claims: 3
CLMN
       Exemplary Claim: 1
ECL
       23 Drawing Figure(s); 23 Drawing Page(s)
DRWN
LN.CNT 1993
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       VH3 and VH4 type immunoglobulins display superantigen-type
AB
       binding affinity for the HIV gp120 envelope glycoprotein. VH3 and VH4
       type antibody molecules, including IgG and IgM, are shown to
       suppress HIV infection in vivo and in vitro. Determining the level of
       such antibody molecules is correlated to the advancement of
       HIV disease state.
L10 ANSWER 25 OF 35 USPATFULL
       97:99175 USPATFULL
ΑN
       Recombinant antibodies for human therapy
ΤI
       Newman, Roland A., San Diego, CA, United States
IN
       Hanna, Nabil, Olivenhain, CA, United States
       Raab, Ronald W., San Diego, CA, United States
       IDEC Pharmaceuticals Corporation, San Diego, CA, United States (U.S.
PA
       corporation)
       US 5681722 19971028
PΙ
       US 1995-478039 19950607 (8)
ΑT
       Division of Ser. No. US 1995-379072, filed on 25 Jan 1995 which is a
RLI
       continuation of Ser. No. US 1992-912292, filed on 10 Jul 1992, now
       abandoned which is a continuation-in-part of Ser. No. US 1992-856281,
       filed on 23 Mar 1992, now abandoned which is a continuation-in-part of
       Ser. No. US 1991-735064, filed on 25 Jul 1991, now abandoned
DT
       Utility
       Primary Examiner: Feisee, Lila; Assistant Examiner: Bansal, Geetha P.
EXNAM
       Burns, Doane, Swecker & Mathis, LLP
LREP
       Number of Claims: 8
CLMN
ECL
       Exemplary Claim: 1
       33 Drawing Figure(s); 26 Drawing Page(s)
DRWN
LN.CNT 2117
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Chimeric antibodies including an Old World monkey portion and
AB
       a human portion, nucleic acid encoding such antibodies, Old
       World monkey monoclonal antibodies, and methods for their
       production and use.
L10 ANSWER 26 OF 35 USPATFULL
       97:73287 USPATFULL
AN
       Recombinant antibodies for human therapy
ΤI
       Newman, Roland A., San Diego, CA, United States
IN
       Hanna, Nabil, Olivenhain, CA, United States
       Raab, Ronald W., San Diego, CA, United States
       Idec Pharmaceuticals Corporation, San Diego, CA, United States (U.S.
PA
       corporation)
       US 5658570
                  19970819
ΡI
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Continuation of Ser. No. US 1992-912292, filed on 10 Jul 1992, now

US 1995-379072 19950125 (8)

ΑI

RLI

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abandoned which is a continuation-in-part of Ser. No. US 1992-856281,
       filed on 23 Mar 1992, now abandoned which is a continuation-in-part of
       Ser. No. US 1991-735064, filed on 25 Jul 1991, now abandoned
DT
       Utility
EXNAM
      Primary Examiner: Feisee, Lila
       Burns, Doane, Swecker & Mathis
CLMN
      Number of Claims: 38
ECL
       Exemplary Claim: 1
       26 Drawing Figure(s); 26 Drawing Page(s)
DRWN
LN.CNT 1829
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Chimeric antibodies including an Old World monkey portion and
       a human portion, nucleic acid encoding such antibodies, Old
      World monkey monoclonal antibodies, and methods for their
       production and use.
    ANSWER 27 OF 35 USPATFULL
L10
       97:70717 USPATFULL
ΑN
       Oral vaccine comprising anti-idiotypic antibody to
TI
       chlamydia glycolipid exoantigen and process
       MacDonald, Alex Bruce, Hatfield, MA, United States
TN
       Whittum-Hudson, Judith A., Elkton, MD, United States
       Saltzman, William Mark, Baltimore, MD, United States
       The Johns Hopkins University, Baltimore, MD, United States (U.S.
PA
       corporation)
       University of Massachusetts, Amherst, MA, United States (U.S.
       corporation)
       us 5656271 19970812
PΙ
       US 1995-466752 19950606 (8)
ΑI
       Continuation of Ser. No. US 1994-213863, filed on 16 Mar 1994, now
RLI
       abandoned which is a continuation-in-part of Ser. No. US 1993-34572,
       filed on 19 Mar 1993
       Utility
DΤ
EXNAM Primary Examiner: Loring, Susan A.
CLMN
       Number of Claims: 15
       Exemplary Claim: 1
ECL
       19 Drawing Figure(s); 10 Drawing Page(s)
DRWN
LN.CNT 2188
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       A genus specific chlamydia oral or injectable vaccine is provided which
       comprises an anti-idiotype antibody capable of
       producing in an animal an anti-idiotypic antibody or
       Fab fragment thereof enclosed in microspheres formed of a
       pharmacologically acceptable polymer is capable of producing in an
       animal an anti-anti-idiotypic immune response (serum
     antibody, secretory antibody or T-cell responsee)
       which recognizes a glycolipid exoantigen (GLXA) of chlamydia. The oral
       or injectable vaccine is produced from an idiotypic
     antibody to GLXA which, in turn, is utilized to produce the
       anti-idiotypic antibody.
L10 ANSWER 28 OF 35 EMBASE COPYRIGHT 2000 ELSEVIER SCI. B.V.
     97316363 EMBASE
AN
     1997316363
DN
     Idiotypic vaccination in B-cell malignancies.
ΤI
     Bianchi A.; Massaia M.
ΑU
     Dr. A. Bianchi, Div. Universitaria di Ematologia, DMOS, AOSGBT, Via
CS
Genova
     3, 10126 Torino, Italy. maxmass@iol.it
    (Molecular Medicine Today, (1997) 3/10 (435-441).
     Refs: 33
     ISSN: 1357-4310 CODEN: MMTOFK
PUI S 1357-4310(97)01105-2
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United Kingdom

CY

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FS
     016
             Cancer
     025
             Hematology
     026
             Immunology, Serology and Transplantation
     037
             Drug Literature Index
LΑ
     English
     English
SL
    CImmunoglobulins contain unique portions, collectively termed
     idiotypes, that can be recognized by the immune system.
     Idiotypes expressed by tumor cells in B-cell malignancies can be
   regarded as tumor-specific antigens and targets for vaccine
immunotherapy.
     Haptens and adjuvants, including cytokines, have been used in several
     animal models to increase idiotype immunogenicity and establish
     protective anti-idiotype immunity. These results have been
     extended by the use of DNA technology, and this has led to the
development
     of a new generation of immunogens, namely fusion proteins and naked-DNA
     (vaccines. The central role of antigen-presenting cells as
     initiators of anti-idiotype immune responses has also been
     recognized. Guided by the experimental data, idiotypic
    vaccination has come into medical use in patients with lymphoma and
     multiple myeloma.
L10 ANSWER 29 OF 35 USPATFULL
       96:116114 USPATFULL
ΑN
       Induction of cytotoxic T-lymphocyte responses
ΤI
       Raychaudhuri, Syamal, San Diego, CA, United States
IN
       Rastetter, William H., Rancho Santa Fe, CA, United States
       IDEC Pharmaceutical Corporation, San Diego, CA, United States (U.S.
PA
       corporation)
       US 5585103 19961217
PI
       US 1992-919787 19920724 (7)
ΑI
       Continuation-in-part of Ser. No. US 1991-735069, filed on 25 Jul 1991,
RLI
       now abandoned
DT
       Utility
       Primary Examiner: Mosher, Mary E.
EXNAM
       Burns, Doane, Swecker & Mathis, LLP
       Number of Claims: 20
CLMN
ECL
       Exemplary Claim: 1
       14 Drawing Figure(s); 9 Drawing Page(s)
LN.CNT 1139
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Methods and compositions useful for inducing a cytotoxic T lymphocyte
       response (CTL) in a human or domesticated or agriculturally important
       animal. The method includes the steps of providing the antigen to which
       the CTL response is desired and providing an antigen formulation which
       comprises, consists, or consists essentially of two or more of a
       stabilizing detergent, a micelle-forming agent, and an oil. This
antigen
       formulation is preferably lacking in an immunostimulating peptide
       component, or has sufficiently low levels of such a component that the
       desired CTL response is not diminished. This formulation is provided as
       a stable oil-in-water emulsion.
L10 ANSWER 30 OF 35 USPATFULL
       95:38596 USPATFULL
AN
       Monoclonal antibody L53 which recognizes a human
ΤI
       tumor-associated antigen
       Hellstrom, Ingegerd, Seattle, WA, United States
IN
       Hellstrom, Karl E., Seattle, WA, United States
       Marquardt, Hans, Mercer Island, WA, United States
       Johnston, Janet, Seattle, WA, United States
       Oncogen Limited Partnership, United States (U.S. corporation)
PΑ
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DT

Journal; General Review

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US 1993-20256 19930218 (8)
ΑI
       Continuation of Ser. No. US 1990-533371, filed on 5 Jun 1990, now
RLI
       abandoned
DΤ
       Utility
       Primary Examiner: Hutzell, Paula K.
EXNAM
       Merchant, Gould, Smith, Edell, Welter & Schmidt
LREP
       Number of Claims: 13
CLMN
       Exemplary Claim: 1
\mathsf{ECL}
       No Drawings
DRWN
LN.CNT 1146
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is concerned with novel monoclonal
     antibody L53 which binds strongly to a glycoprotein antigen
       associated with human tumors, including carcinomas of the colon,
breast,
       and lung, as well as melanomas. The antibody binds to normal
       human cells to a much lesser degree than to tumor cells. The
     antibody finds use in diagnostic methods for as the detection of
       malignant cells associated with tumors. Also disclosed is a novel
       70,000-75,000 dalton glycoprotein antigen recognized by MAb L53. The
L53
       antigen is found on the cell surface of human tumor cells. The amino
       terminal amino acid sequence of this antigen is: ##STR1## in which X
       represents an unidentified amino acid.
    ANSWER 31 OF 35 MEDLINE
L10
     96135362
                  MEDLINE
ΑN
     96135362
DN
     A genetic approach to idiotypic vaccination for B
ΤI
    cell lymphoma.
     Stevenson F K; Zhu D; King C A; Ashworth L J; Kumar S; Thompsett A;
ΑU
     Hawkins R E
     Molecular Immunology Group, Tenovus Laboratory, Southampton University
CS
     Hospitals, United Kingdom.
     ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1995 Nov-27) 772 212-26.
SO
     Ref: 23
     Journal code: 5NM. ISSN: 0077-8923.
CY
     United States
     Journal; Article; (JOURNAL ARTICLE)
DT
     General Review; (REVIEW)
     (REVIEW, TUTORIAL)
     English
LΑ
     Priority Journals; Cancer Journals
FS
ΕM
     199604
     Idiotypic immunoglobulin expressed by a B cell tumor presents a
AB
   Clear tumor antigen which could be attacked by vaccination of the host.
     Vaccination with idiotypic protein has been shown to induce
    (protective immunity against lymphoma, but application to patients is
     limited by the requirement of "personal" vaccines for each
     patient. A genetic approach enables V-region sequences encoding
     idlotypic antigen to be rescued from tumor biopsies, and to be
     assembled as scFv fragments. These can be expressed in bacteria to
prodúce
     recombinant protein, or used directly as naked DNA vaccines.
     Intramuscular injection of idiotypic DNA from a mouse B
     cell lymphoma induces low levels of syngeneic anti-
     idiotypic antibody in serum. Response can be stimulated
     by co-injection of DNA plasmids encoding either IL-2 or GM-CSF, and T
     cells which proliferate in response to idiotypic IgM are
     generated. However, protection against tumor appears to be blocked by
     continuing secretion of idiotypic antigen from the persisting
     vaccine vector, which forms immune complexes with serum antibody
     . Methods for regulating the level of scFv to engage the immune system,
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PΙ

US 5411884 19950502

but not to block the effector arm are being investigated. Similar control will be applicable to the cytokine vectors, which can deliver encoded cytokines designed to activate immune pathways for tumor destruction. Experience gained in lymphoma may be extended to other tumors with tumor antigens. ANSWER 32 OF 35 USPATFULL L10 93:56702 USPATFULL ΑN Anti-idiotype antibodies reactive with shared TIidiotopes expressed by B cell lymphomas and autoantibodies Miller, Richard A., 8 Ohlone, Portola Valley, CA, United States 94025 IN US_5227159 19930713-7 PΙ US 1992-898246 19920612 (7) ΑI Continuation of Ser. No. US 1990-467405, filed on 22 Jan 1990, now RLI abandoned which is a continuation-in-part of Ser. No. US 1989-304745, filed on 31 Jan 1989, now abandoned Utility DTPrimary Examiner: Lacey, David L.; Assistant Examiner: Budens, Robert EXNAM D. Woolcott, Kenneth J.; Burgoon, Jr., Richard P. LREP Number of Claims: 12 CLMN Exemplary Claim: 1 ECL 10 Drawing Figure(s); 6 Drawing Page(s) DRWN LN.CNT 1351 CAS INDEXING IS AVAILABLE FOR THIS PATENT. B-cell lymphomas express surface immunoglobulin (immunoglobulin) ΑB containing unique idiotypic (idiotype) determinants which may be exploited as tumor specific markers. The inventor has produced murine monoclonal antibodies (MAbs) reactive with the idiotype marker derived from 67 patients with low grade, follicular, small cleaved cell lymphoma. Out of 199 monoclonal antibodies, 47 (24%) were found to react with pooled normal human serum immunoglobulin in concentrations ranging from 0.6 .mu.g/ml to 160 .mu.g/ml. Of these 40 monoclonal antibodies, 90% cross-reacted with idiotype present in normal serum in levels <50 .mu.g/ml. Thirty-two of these anti-idiotypes were directed against a shared idiotope expressed on another patient's lymphoma cells. The frequency of shared idiotope expression defined by each antibody ranged from 0.26% to 3.9% of the B-cell lymphomas tested. A panel of five anti-idiotype antibodies reacted with 80% of AIDS associated lymphomas. Based on the reactivity with these monoclonal antibodies, tumors could be grouped into distinct families. In aggregate, these 32 monoclonal antibodies reacted with a total of 108 of 332 B cell lymphoma cases (32.5%), including 35 of 116 follicular, small cleaved cell lymphomas (30%). Many of these anti-shared idiotopes reacted with more than one histopathologic subtype of lymphoma. Antiidiotypes have been used in B-cell lymphoma diagnosis and therapy. Moreover, applicant has discovered at least seven anti-shared idiotype antibodies that cross react with autoantibodies, e.g., 16.6 and RF. The development of a library of anti-idiotypes reactive with shared idiotopes should facilitate these clinical studies by obviating the need to develop a customized hybridoma for each patient. L10 ANSWER 33 OF 35 MEDLINE

- AN 93226047 MEDLINE
- DN 93226047
- Idiotype/granulocyte-macrophage colony-stimulating factor fusion protein as a vaccine for **B=cell lymphoma** [see comments].
- CM Comment in: Nature 1993 Apr 22;362(6422):695

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Department of Medicine, School of Medicine, Stanford University,
CS
     California 94305.
    SO
     Journal code: NSC. ISSN: 0028-0836.
CY
     ENGLAND: United Kingdom
     Journal; Article; (JOURNAL ARTICLE)
DT
LΑ
     Priority Journals; Cancer Journals
FS
EΜ
     199307
     To produce a vaccine against cancer, antigens must be found that are
AB
     preferentially expressed by tumour cells and can induce an immune
     against the tumour. The variable regions of the immunoglobulin modecules expressed on malignant B.cells. idiotypes are tumour-specific,
     but are weak immunogens. To induce an immune response in animals or
     humans, the idiotypic protein has therefore to be chemically
     coupled to a strongly immunogenic protein and mixed with an adjuvant. The
     resulting response can protect animals from subsequent tumour challenge,
     and cure animals with established tumours in combination with
     chemotherapy. Gramulocyte-macrophage colony-stimulating factor (GM-CSF)
     automents antigen presentation in a variety of cells. Here we show that by
     fusting a tumour-derived idiotype to GM-GSF, it can be converted
     interest trong immunogen capable of inducing idiotype-specific)
     antibodies without other carrier proteins or adjuvants and of
     protecting recipient animals from challenge with an otherwise lethal dose
     of tumour cells. This approach may be applicable to the design of
     vaccines for a variety of other diseases.
    ANSWER 34 OF 35 USPATFULL
L10
       92:102981 USPATFULL
ΑN
       Monoclonal antibody to novel antigen associated with human
ΤI
       Hellstrom, Ingegerd, Seattle, WA, United States
TN
       Hellstrom, Karl E., Seattle, WA, United States
       Marquardt, Hans, Mercer Island, WA, United States
       Oncogen, Seattle, WA, United States (U.S. corporation)
PA
       US 5171665 19921215
PΙ
ΑI
       US 1989-339142 19890417 (7)
       Utility
DT
       Primary Examiner: Kepplinger, Esther L.; Assistant Examiner: Scheiner,
EXNAM
       Toni R.
       Mandel, SaralynnSheldon & Mak
LREP
       Number of Claims: 24
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 1173
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is concerned with a novel monoclonal
AB
     antibody which binds strongly to a protein antigen associated
       with human tumors, including carcinomas of the colon and lung. The
     antibody binds to normal human cells to a much lesser degree
       than to tumor cells. The antibody finds use both in diagnostic
       methods such as the detection of malignant cells associated with tumors
       and in therapeutic methods for treatment of humans with tumors. Also
       disclosed is a novel 66,000 dalton glycoprotein antigen found on the
       cell surface of human tumor cells. The amino terminal amino acid
       sequence of this antigen is: ##STR1## in which X represents an
       unidentified amino acid.
L10 ANSWER 35 OF 35 USPATFULL
ΑN
       92:61852 USPATFULL
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Monoclonal antibody to novel antigen associated with human

Comment in: Nature 1993 Aug 5;364(6437):493

ΑU

ΤI

Tao M H; Levy R

Hellstrom, Karl E., Seattle, WA, United States IN Hellstrom, Ingegerd, Seattle, WA, United States Marquardt, Hans, Mercer Island, WA, United States Yoneyama, Yoshitaka, Bellevue, WA, United States Oncogen Limited Partnership, Seattle, WA, United States (U.S. PΑ corporation) PΙ US 5134075 19920728 US 1989-312640 19890217 (7) ΑI DTUtility Primary Examiner: Kepplinger, Esther L.; Assistant Examiner: Bidwell, EXNAM Carol E. Mandel, SaraLynn LREP Number of Claims: 21 CLMN Exemplary Claim: 8 ECLNo Drawings DRWN LN.CNT 1097 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention is concerned with a novel monoclonal AB antibody which binds strongly to a protein antigen associated with human tumors, including carcinomas of the colon, breast, ovary and lung, as well as melanomas and sarcomas. The antibody binds to normal human cells to a much lesser degree than to tumor cells. The antibody finds use both in diagnostic methods such as the detection of malignant cells associated with tumors and in therapeutic methods for treatment of humans with tumors. Also disclosed is a novel

100,000 dalton glycoprotein antigen found on the cell surface of human tumor cells. The amino terminal amino acid sequence of this antigen is:

##STR1## in which X represents an unidentified amino acid.

=> s B cell lymphoma vaccine!

L12

L11 0 B CELL LYMPHOMA VACCINE!

=> s vaccine comprising immunoglobulin!

O VACCINE COMPRISING IMMUNOGLOBULIN!

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BELLSTEIN adds new search [ields
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MEDLINE SDI run of October 8, 2002
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PHARMAML offering one free connect hour in February 2003
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435/172.2; 435/70.21
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COMMAND INTERRUPED
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If this message appears repeatedly, please notify the Help Desk.
Enter "HELP STN" for information on contacting the nearest STN Help
Desk by telephone or via SEND in the STNMAIL file.
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INCLS: 530/388.250, 530/389.300, 435/240.270
NCLM: 530/387.300
NCLS: 530/388.250, 530/389.300
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1515 L1 AND B(A) CELL
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387 L2 AND IDIOTYP?
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2 FILES SEARCHED...
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Methods for producing high affinity anti-human IgB-monoclonal antibodies which binds to IgE on IgEabearing B cells but not
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INCLS: 530/388.250; 435/240.270; 435/172.200; 435/070.210; 435/240.200
NCLM: 530/387.300
NCLS: 435/070.210; 530/388.250
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Chang, Tse-wen, Houston, TX, United States
Tanox Blosystems, Inc., Houston, TX, United States (U.S. corporation)
US 1422258
US 1988-226421
19880729 (7)
Continuation-in-part of Ser. No. US 1987-140036, filed on 31 Dec 1987, now abandoned
Utility
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Continuation of Ser. No. US 1988-291068, filled on 28 Dec 1988, now abandoned which is a continuation-in-part of Ser. No. US 1988-226421, filled on 29 Jul 1988 which is a continuation-in-part of Ser. No. US 1987-140036, filed on 31 Dec 1987, now abandoned
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ICS: C12N005-20, C07K016-42
530/387; 530/388.25; 435/240.27; 435/172.2; 435/240.27; 435/70.21
INDEXING IS AVAILABLE FOR THIS PATENT.
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ICS: CL2NOOS-10; CL2NOOS-20
530/387.3; 530/388.25; 435/240.2; 435/240.27; 435/172.2; 435/70..
INDEXING IS AVAILABLE FOR THIS PATENT.
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INCLS: 530/387.300; 530/388.250; 530/388.730; 435/240.270
NCLM: 530/387.300; 530/388.250; 530/388.730
NCLS: 530/387.300; 530/388.250;
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Sun, Bill N., Houston, TX, United States
Sun, Cecily R., Houston, TX, United States
Sun, Susystems, Inc., Houston, TX, United States (U.S.,
US 542021
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Anti-idiotype antibodies specific for the parotope
antibodies which bind to IgE-bearing B cells but no
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INCLS: 530/388.250, 435/240.270; 435/070.210
NCLM: 435/452.000
NCLS: 435/070.210; 530/388.250
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95:50082 USPATFULL
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AGAINST SOFT WE USPATITUDE.

Starburst conjugates
Fapturst conjugates
Tomalia, Donald A., Midland, MI, United States
Kaplan, Donald A., Midland, MI, United States
Kraptan, Donald A., Midland, MI, United States
Kraptan, Roberta C., Midland, MI, United States
Tomilnson, Ian A., Midland, MI, United States
Fazio, Michael J., Midland, MI, United States
Fazio, Michael J., Midland, MI, United States
Fazio, Michael J., Midland, MI, United States
Milson, Larry R., Beaverton, MI, United States
Wilson, Larry R., Beaverton, MI, United States
The Dow Chemical Company, Midland, MI, United States
The Dow Chemical Company, Midland, MI, United States
The One Continuation in-part of Ser. No. US 1980-1864 Ser. No. US 1987-87266, filed on 18 Aug 1987, now abandoned which is a continuation-in-part of
Ser. No. US 1986-897455, filed on 18 Aug 1986, now abandoned
Utility

Granted LS ANSWER 6 OF 40 USPATFULL

AN 93:22794 USPATFULL

IT Molecular recognition units

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N Wolecular Thoma J. Yardley, PA, United States

Alvarez, Vernon L., Morrisville, PA, United States

Alvarez, Vernon L., Morrisville, PA, United States

Radcliffe, Robert D., Tiusville, PA, United States

Radcliffe, Robert D., Tiusville, PA, United States

Radcliffe, Robert D., Tiusville, PA, United States

I Styden Corporation, Princeton, NJ, United States

US 1990-519702

I Continuation-in-part of Ser. No. US 1988-291730, filed on 29 Dec 1988, United States

United States 424/009.364; 424/078.170; 514/772.300; INCLM: 424/001.490
INCLS: 424/001.00; 424/078.080; 424/078.100; 424/009.000; 424/078.170;
424/001.370; 424/001.530; 424/001.690; 424/001.650; 521/025.000;
521/028.000; 436/173.000; 436/806.000; 514/772.100; 514/772.300 INCLM: 530/324.000 INCLS: 530/326.000; 424/001.100; 424/002.000; 436/545.000; 436/546.000 NCLM: 530/324.000 NCLS: 436/545.000; 436/546.000; 530/326.000 [5]
ICM: A61K043-00
ICS: A61K031-785; A61K031-80
ICS: A62K031-74; A61K031-785; A61K031-80
424/9.83; 424/78.13; 424/78.13; 424/63; 424/63; 424/647; 424/648; 424/647; 424/648; 424/647; 424/648; 424/617; 424/64; 521/28; 521/25; 436/173; 436/806; 514/772.1; 514/772.3
INDEXING IS AVAILABLE FOR THIS PATENT. ICS: C12N005-20 530/387; 530/387.2; 530/387.3; 530/388.25; 530/388.73; 435/240.27 INDEXING IS AVAILABLE FOR THIS PATENT. 424/001.530; 424/001.650; 424/001.690; 424/009.360; 424/009.400, 434/09.400; 434/09.360; 424/078.080; 424/078.100; 434/078.370; 436/173.000; 436/806.000; 514/772.100; 521/025.000; 521/028.000 [5] ICM: CO7K007-08 ICS: CO7K007-10; A61K043-00; G01N033-533 436/513; 436/545; 436/546; 530/324; 530/326; 424/2 USPATFULL OF 40 NCLM: NCLS: DT CN CNT NINCL CNT DT FS LN.CN INCL NCL CAS PA PI AI RLI PA PI AI RLI EXF NCL I I A F S SALI

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CAS INDEXING IS AVAILABLE FOR THIS PATENT

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436/809.000;
436/808.000;
530/326.000;
530/387.000
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436/543.000;
530/324.000;
530/329.000;
530/868.000
ANSWER 7 OF 40 USPATFULL
91:96281 USPATFULL
Anti-idiotype antibodies induced by synthetic polypeptides
Carson, Dennis A., Del Mar, CA., United States
Carson, Dennis A., Del Mar, CA., United States
Composition of the Carson of 
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    INCLM: 424/085.910 .
INCLS: 530/387.000
NCLM: 424/183.100
NCLS: 424/085.000; 424/809.000; 530/387.200; 530/388.730; 530/391.700; 530/862.000; 530/864.000; 530/866.000
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Whr. Jonathan W. Dallas, TX, United States
Uhr. Jonathan W. Dallas, TX, United States
Board of Regents, The University of Texas System, Austin, TX, United
States (U.S. corporation)
US 4792447
US 4792447
US 1983-498754
Continuation in part of Ser. No. US 1981-286090, filed on 23 Jul 1981,
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LINCLM: 435/007.920
LINCLM: 435/10.000; 435/007.930; 435/965.000; 424/088.000; 436/424.000; 436/539.000; 436/543.000; 436/543.000; 436/436/4000; 436/543.000; 436/436/4000; 436/43.000; 436/43.000; 436/43.000; 436/43.000; 530/325.000; 530/325.000; 530/325.000; 530/327.000; 530/327.000; 530/320.000; 530/330.000; 530/320.000; 530/330.000; 530/320.000; 434/197.110; 424/197.110; 424/197.110; 424/197.110; 424/197.110; 424/197.110; 424/197.110; 424/197.110; 436/47.000; 436/518.000; 436/539.000; 436/518.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.0000; 530/330.000; 530/330.000; 530/330.000; 530/330.000; 530/330.000
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ICM: A61K039-395
ICS: A61K039-44; C07K015-00; C07K017-00
424/85; 424/92; 424/87; 260/112R; 435/172.2; 530/387
INDEXING IS AVAILABLE FOR THIS PATENT.
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ICM: GOIN033-53
ICS: GOIN033-543; C07K015-14; C07K007-00
435/810; 436/808
INDEXING IS AVAILABLE FOR THIS PATENT
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                ANSWER 7
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1995024221 PCTFULL ED 20020514
BIOACTIVE AND/OR TARGETED DENDRIMER CONJUGATES
CONJUGUES DENDRIMERES BIOACTIFS ET/OU CIBLES
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THE DOW CHENICAL COMPANY;
THE DOW CHENICAL COMPANY;
BENDRITECH INCORPORATED;
REGENTS OF THE UNIVERSITY OF MICHIGAN;
TOMALIA, Donald, A.;
BAKER, James, N.;
BENTHERS, Herbert, M., II;
CHENO, Roberta, C.;
FAZIO, Michael, J.,;
HEDSTRAND, David, M.;
JOHNSON, Jennifer, A.;
KAPLAN, Donald, A.;
KAPLAN, Donald, A.;
KARAMP, SCOTT, L.;
KAUPER, WILliam, J., Jr;
KUKOWSKA-LATALLO, Jolanta;
MAXON, Bartley, D.;
PIEHLER, Lars, T.;
TOMLINSON, Larry, R.;
WILLON, Larry, R.;
WILLON, Larry, R.;
                                                      TOWALIA, Donald, A.;
BAKER, James, R.;
BIELINSKA, Anna, U.;
BROTHERS, Herbert, M., II;
FAZIO, Michael, J.;
FAZIO, Michael, J.;
HEDSTRAND, David, M.;
JOHNSON, Jennifer, A.;
KAPLAN, Donald, A.;
KAPLAN, Donald, A.;
KARDER, William, J., Jr;
KUKOWSKA-LATALLO, Jolanta;
MAXON, Bartley, D.;
TOWALNSON, Larry, R.;
WILSON, Larry, R.;
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INCLS: 530/387.000; 530/388.000; 424/085.000; 436/547.000; 436/548.000;
NCLM: 424/131.100
NCLS: 435/070.300; 435/070.400; 436/547.000; 436/548.000; 530/387.200; 530/389.400; 530/863.000; 530/865.000
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1995029700 PCTFULL ED 20020514
SYNTHETIC VACCINE FOR PROTECTION AGAINST HUMAN IMMUNODEFICIENCY VIRUS
INFECTION
VACCIN DE SYNTHESE PROTECEANT CONTRE L'INFECTION PAR LE VIH
HAYNES, Barton, F.;
PALKER, Thomas, J.
DUE UNIVERSITY
ENGLISH
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W: MO 1995-US5465
AU CA JP AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE NO 1995-US5465
A 19950428
A61K039-21
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CA JP US AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE
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35,404 19940429
Scripps Clinic and Research Foundation, La Jolla, CA, United States (U.S. corporation) 19870728 US 1984-614102 19840524 (6) Utility Granted
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                                                                                                                                                                                                                                                                                                                  ANSWER 10 OF 40 PCTFULL COPYRIGHT 2003 Univentio DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        VON FELDT, Joan, M.;
KIBBER-EMMONS, Thomas;
WEINER, David, B.;
WILLIAMS, WIlliam, V.
THE TRUSTEES OF THE UNIVERSITY OF PENNSYLVANIA;
THE WISTAR INSTITUTE;
VON FELDT, Joan, M.;
KIBER-EMMONS, Thomas;
WEINER, David, B.;
WILLIAMS, William, V.
                                                                                                                                                                                                                          IC (4)
ICM: CO7K003-08
EXF 260/112B; 424/8S; 436/547; 435/68
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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PRETARGETING METHODS AND COMPOUNDS PROCEDES ET COMPOSES DE PRECIBLAGE MEYER, Damon, L.;
MALLETT, ROBEET, ROBEE
M: 524221 A1 19950914
W: AU BR CA CN CZ EE F1 GE HU
SK UA US US US US US US US
LU MC NL PT SE
WO 1995-US3045 A 19950307
US 1994-8/207,494 19940930
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AALKO47-48
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AXWORTHY, Donald, B.;
GUSTAVSON, Linda, M.
NEORX CORPORATION
English
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A61K038-00
A61K038-03; A61K038-16; C07K001-00; C07K007-00; C07K007-04; C07K007-08; C07K011-00; C07K014-00; C07K019-00; C07K016-24; C07K016-42; C12P021-00; C12P021-02; C12P021-04

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ANSWER 13 OF 40

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TREATMENT OF AUTOIMNUE DISEASE
TRAITEMENT DE MALADIES AUTO-IMMUNES
BERNARD, Claude, Charles, Andre;
KERLERO DE ROSBO, Nicole, Claude, Marie
LA TROBE UNIVERSITY;
BERNARD, Claude, Charles, Andre;
KERLERO DE ROSBO, Nicole, Claude, Marie
ENRARD, Claude, Charles, Andre;
BERNARD, Claude, Charles, Andre;
KELLERO DE ROSBO, Nicole, Claude, Marie
English
We 9507096
WH BE BG BR BY CA CH CN CZ DE DK EE ES SD WW ANSWER 16 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1995015180 PCFFULL ED 20020514
ANTIGODIES THAT MINT ACTIONS OF NEUROTROPHINS
ANTICORPS IMITANT LES EFFETS DES NEUROTROPHINES COPYRIGHT 2003 Univentio II C S M R ¥ W ₩ CZ DE I MD MG N KE MW S WO 9515770 A1 19950615 W: CA JP AT BE CH DE DK ES FR GB GR IE WW 1994-US14223 A 19941209 US 1993-8/164,302 19931209 MG I US 1993-8/164,302 19931209 A61K051-00; A61K051-10; A61K049-00; A61K047-48 Z Q Z CLARY, Douglas, 0.;
WESKAM, Cisela,
AUSTIN, Leeann;
REICHARDT, Louis, F.
FIEREGENTS OF THE UNIVERSITY OF CALIFORNIA
English MO 9515180

AM AT AU BB BG RB BY CA CH CN
KE KG KR KE LK LR LT LU LV
RU SD SE SI SK TJ TT UN UZ VN
FR GB GR IE IT LU NC NL PT SE
WO 1994-US13708

AUS 1993-8/162.597

ASTRONO1-38; GOIN033-53 SAES S S S A1 19950316 U BB BG BR BY CA CH C P KR KZ LK LR LT LU L E SI SK TJ TT UA US U Al 19950615 CA JP AT BE CH DE DK ES FR 4174 A 19941207 63,188 19931207 ANSWER 15 OF 40 PCTFULL COPYRIGHT
1995015770 PCTFULL ED 20020514
PRETARGETING METHODS AND COMPOUNDS
PROCEDES ET COMPOSES DE PRECIBLAGE
GRAVES, SCOCT, S.;
BLORN, Michael, J.;
RENO, John, M.;
AXWORTHY, Donald, B.;
FRIYZBERG, Alan, R.;
FREDORE, Louis, J. AU KP WO 1994-US14174 US 1993-8/163,188 AT KG SD KE A Patent WO 9515979 Patent WO 9515180 W: English Patent DT PI DS AI PRAI ICM ICS LS AN TIEN TIFR IN PA LA DT PI DS AI PRAI ICS LS AN TIEN TIFR IN LS AN TIEN TIFR IN AI PRAI ICM ICS PA DT PI DS ΡA PI DI

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199500431 PCTFULL ED 20020514
METAL RADIOWUCIDE LABELED PROTEINS FOR DIAGNOSIS AND THERAPY
PROTEINES MARQUEES ANCE DES RADIOWUCLEIDES METALLIQUES DESTINEES A UNE
PRITIZERATION DIAGNOSTIQUE ET THERAPEUTIQUE
KASINA, Sudhakar;
GUSTRANOW, Linda, M.
MEDRE COPPERATION
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KAY, William, W.;

COLLINGON, S., Karen;

CLOUTHIER, Sharon, C.

UNIVERSITY OF VICTORIA INNOVATION AND DEVELOPMEN T CORPORATION;
                                                                                                                           BHAT, Neelima, M.;
BIEBER, Marcia, M.;
TENG, Nelson, N., H.
THE BOARD OF TRUSTEES OF THE LELAND STANFORD JUNIOR UNIVERSITY;
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                                                                                   ANSWER 18 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1995003770 PCTFULL B 200205114 METHODS FOR B-CELL POPULATION CONTROL PROCEDES DE REGULATION DE LA POPULATION DES LYMPHOCYTES
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M. AM AT AU BB BG BR BY CA CH CN CZ DE DR KG KR KZ LK LT LU LV MD MG MN MW NI SE SI SK TJ TI UA US UZ VN KE MW SD AI GR IE IT LU MC NL PT SE BF BJ CF CG CI TD TG WO 1994-US8793

WO 1994-US8793

A 19940802

A613001-00

A61K038-16; A61K039-00; C07K016-30; C07K001-00
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BIEBER, Marcia, M.;
TENG, Nelson, N., H.
                     WO 1994-AU522
AU 1993-PM 1030
A61K038-17
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English
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WO 9503770
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PEPTIDES IMMUNOTHERAPEUTIQUES DERIVES DE LA TOXINE-1 DU SYNDROME DU CHOC TOXIQUE, ANTICORPS CONTRE CELLE-CI, LEURS UTILISATIONS DANS DES COMPOSITIONS PHARMACEUTIQUES ET EN DIAGNOSTIC KP KR KZ TT UA US SE BF BJ JP KG KP SK TJ TT NL PT SE ES FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML MR NE SN TD TG A 19940426 ANSWER 23 OF 40 PCTFULL COPYRIGHT 2003 Univentio
19940A164 PCTFULL ED 20020513
HUMAN MONOCLONAL ANTIBODIES AND PROCESSES AND MATERIALS FOR MAKING SUCH
ANTIBODIES
ANTICORPS MONOCLONAUX HUMAINS, ET PROCEDES ET MATERIAUX DE FABRICATION ANSWER 22 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1994025483 PCTFULL ED 20020513 IMMUNOTHERAPEUTIC PEPTIDES DERIVED FROM TOXIC SHOCK SYNDROME TOXIN-1, AMTIBODIES THERETO, THEIR USES IN PHARMACEUTICAL COMPOSITIONS AND DIAGNOSIS ANIMAUX TRANSGENIQUES CAPABLES DE PRODUIRE DES ANTICORPS HETEROLOGUES ANSWER 21 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1994025585 PCTFULL ED 20020513 TRANSCENIC NON-HUMAN ANIMALS CAPABLE OF PRODUCING HETEROLOGOUS ANTIBODIES JP SK PT SE MC FI GB HU S SD SE SI S LU MC NL E FI GB GE H RO RU SD S I IE IT LU M CL2N015-31 C1220001-68; G01N033-569; C07K013-00; C12P021-08; C12Q001-04; C12N005-12 CO7KO13-00; C12PO21-08; A61KO37-02; C07KO07-10; C07KO07-06; A61KO39-40; G01NO33-50; G01NO33-68; A61KO39-39 ES IT 3 ES 23 ES 2 DE DK PT RO GR IE TD TG DE DK NZ PL FR GB NE SN Mils;

Abert, M.

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425585
AT AU BB GB RB BY CA CH CN CZ DE
LK LU LV MG MN MW NL NO NZ PL PT
UZ VN AT BE CH DE DK ES FR GB G
CG CI CM GA GN ML MR NE SN '
WO 1994-US4580
US 1993-8/053, 131
1993-18/155, 301
1993-18/155, 301
1993-1203
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1993-1203
1993-1203 M: 9425483 AT AU BB GG BR BY CA CH CN CZ
KR KZ LK LU LV MG MN MW NL NO
UA US UZ VN AT BE CH DE DK ES
WO 1994-IB140 A 19930503
US 1993-8/058,518 19930503 CZ NO WS CHOW, Anthony, W.; KUM, Winnie, W.; THE UNIVERSITY OF BRITISH COLUMBIA, CHOW, Anthony, W.; KUM, Winnie, W., S. 19930426 KAY, Robert, M.

KAY, Robert, M.

CENHARM INTERNATIONAL, INC.;

LONBERG, Nils;

KAY, Robert, M.

English
Parent
WO 9425585
AT AU BB GG RB BY CK WO 1994-IB205 US 1993-8/054,452 TIFR IN AI PRAI LS AN TIEN L5 AN TIEN LS AN TIEN AI PRAI ICM ICS TIFR AI PRAI ICM ICS ICM ΡA PI DT N ΡA LA DT PI DS

IN PA LA DT DT DS AI CCM ICS LS AN TIFEN IN	
PA LIA DT PI DS AI PRAI ICM	### BHATT, Ramesh, R.; BHATT, Ramesh, R.; AFFYMAX TECHNOLOGIES N.V.; THE BOARD OF TRUSTEES OF THE LELAND STANFORD JR. UNIVERSITY THE BOARD OF TRUSTEES OF THE LELAND STANFORD JR. UNIVERSITY THE BOARD OF TRUSTEES OF THE LELAND STANFORD JR. UNIVERSITY WO 9418345 WO 9418345 A 199400818 WO 1993-8/014,426 US 1993-8/014,426 US 1993-8/155,341 19931115 C12001-68 C12001-60, A61K037-00
LS AN TIEN TIFR IN PA LA	ANSWER 25 OF 40 PCTFUI 1994014469 PCTFUIL ED 3 MULTIVALENT ARBAD ANTICORPS AB1 MULTIVALER RODKEY, L., SCOLT; SFERIAN, PETER, G. BOARD OF REGENTS, THE UN ENGlish
PI DS AI PRAI ICM ICS	WO 9414469 W1 9414469 W2 414469 W3 119940707 W3 41 AU BB BG BR BY CA CH CZ DE DK ES FI GB HU JP KP KR KZ LK LU LV MG MN MM NL NO NZ PL PT RO RU SD SE SK UA UZ VN AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM CA GM ML MR NE SN TD TG W0 1993-US19413 M 19931220 US 1992-7/995,373 M 19931223 A61K037-04 A61K039-385; A61K039-39; CG7K017-02
LS AN TIEN TIFR	ANSWER 26 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1994010294 PCTFULL ED 20020513 HUMAN WARNOCLOMAL ANTIBODIES TO HUMAN PARVOVIRUS AND METHODS OF MAKING AND USING THEREOF ANTICORPS MONOCLOMAUX CONTRE LE PARVOVIRUS CHEZ L'HOMME, PROCEDES D'OBTENTION ET D'UTILISATION

DE CES ANTICORPS

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ANSWER 31 OF 40 PCTFULL COPYRIGHT 2003 Univentio 193201207 PCTFULL ED 20020513

EN MONGCLONAL ANTHODIES WHICH BIND TO SECRETED AND MEMBRANE-BOUND IGE, NOT TO IGE ON BASOPHILS

RANTICORPS MONOCLONAUX QUI SE LIENT A L'IGE SECRETEE ET LIEE AUX MEMBRANES MAIS QUI NE SE LIENT PAS A L'IGE SECRETEE ET LIEE AUX MEMBRANES MAIS QUI NE SE LIENT PAS A L'IGE SECRETEE ET LIEE AUX MEMBRANES MAIS QUI NE SE LIENT PAS A L'IGE SECRETEE ET LIEE AUX GOSSETT, Lani, A.;

SUN, Lee, K.;

SUN, Lee, K.;

SUN, Cecily, R., Y.;

LIOU, Ruey, S.

SUN, Lee, K.;

SUN, Cecily, R.;

SUN, Cecily, R.;

SUN, Ruey, S.
                                                                                                                                                                                                                      ANSWER 30 OF 40 PCTFULL COPYRIGHT 2003 Univentio
1993012227 PCTFULL ED 20020513
TRANSGEBLIC NON-HUMAN ANIMALS CAPABLE OF PRODUCING HETEROLOGOUS
ANTIBODIES
ANTIBODIES
HETEROLOGUES
HETEROLOGUES
HETEROLOGUES
    DE DK ES FI GB HU
RU SD SE SK UA US
NL PT SE BF BJ CF
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RODMAN, TOby, C.
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ZOLLA-PAZNER, Susan;
ARAKELOV, Serguei;
MIROSLAV, Gorny
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ANSWER 35 OF 40 PCTFULL COPYRIGHT 2003 Univentio ANSWER 35 OF 40 PCTFULL ED 20020513 NOLECULAR RECCONSISANCE MOLECULAIRES ROWELL, John, D.; ALVAREZ, Vernon, D.; ALVAREZ, Vernon, L.; RADCLIFFE, Robert, D. CYTOGER CORPORATION English Patent WO 911713 AT BE CA CH DE DK ES FR GB GR IT JP LU NL SE WO 199-103116 A 19910507 A 61702-00 A 617035-14; A61K037-00	ANSWER 36 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1991006305 PCTFULL ED 20020513 OLGCOMERIC IMMUNOCLOBULINE SHURDNOL MALL W.; HARLIS, Linda, J.; RAFF, HOWARG, V. BRISTOL-MYERS SQUIBB COMPANY English Patent WO 9106305 AT AU BE CA CH DE DK ES FI FR GB GR IT JP KR LU NL NO SE WO 1990-USG426 N: 1989-432.700 19891107 A 61K039-40; C12N005-02; C12N015-00	ANSWER 37 OF 40 PCTFULL COPYRIGHT 2003 Univentio 199007713 PCTFULL ED 2002013 MOLECULAR RECCENTITON UNITS UNITES DE RECONNAISSANCE MOLECULAIRE RODWELL, JOH, D.; MCKEARN, Thomas, J. CYTOGEN CORPORATION English Patent WO 9007713 WI 1989-USSB50 A 19891228 WO 1989-291,730 19881229 G C1220001-68; A01N063-00	ANSWER 38 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1990007321 PCTFULL ED 20020513 N HOMING SEQUENCES AND THEIR USES SEQUENCES DE GUIDAGE ET LEURS EMPLOIS WEISSMAN, Irving, L.; FULLMANN, BATK, H. SIEGELAAN, MARK, H. THE BOARD OF TRUSTEES OF THE LELAND STANFORD JUNIOR UNIVERSITY English Patent WO 9007321 A1 19900712
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oligomers, typically of the IgG class, are secreted having two or more immunoglobulin monomers associated together to form tetravalent or hexavalent IgG molecules. The oligomers can be formed by substantially duplicating regions of the light chain, particularly the variable region. Oligomeric antibodies of the IgG isotype cross the placenta and can provide passive immunity to a fetus, which is particularly important for protecting newborns against pathogens such as group B streptococci. ANSWER 36 OF 40 PCTFULL COPYRIGHT 2003 Univentio Novel oligomeric monoclonal antibodies with high avidity for antigen are described. The ANSWER 39 OF 40 PCTFULL COPYRIGHT 2003 Univentio 198906138 PCTFULL ED 20020513 UNIQUE ANTIGENIC EPITOPES ON IGE-BEARING B LYMPHOCYTES D'IMMUNOGLOBULINE E ANSWER 40 OF 40 PCTFULL COPYRIGHT 2003 Univentio 1955002909 PCTFULL ED 20020677 SYNTHETIC POLYPEPTIDES ANTI-IDICATER ANTI-LIDICATER ANTI-LIDICATER INDUITS PAR DES POLYPEPTIDES ANTI-LIDICATER INDUITS PAR DES POLYPEPTIDES A1 19890713 IB CH DE FR GB IT JP KR LJU NL SE SU A 1981729 1981721 19880729 19881116 19881116 A1 19850704 CH DE DK FR GB JP LU NL SE A 19841228 19831228 C12P021-00; A61K047-00; C12N005-00; C12N015-00 SE LU NL MCMILLAN, Seamus, L., H. SCRIPPS CLINIC AND RESEARCH FOUNDATION English ď DE FR GB IT J A 19891110 19881223 19890224 CHANG, TSE-Wen; SUN, Bill, Nai-Chau; SUN, Cecily, Rou-Yun TANOX BIOSYSTEMS, INC. English CARSON, Dennis, A.; HOUGHTEN, Richard; CHEN, Pojen, P.; VAUGHAN, John, H.; LERNER, Richard, A.; AT BE CH AT AU BE AT AU BE WO 1988-US4706 US 1987-140,036 US 1988-226,421 US 1988-229,178 US 1988-2272,243 US 1988-291,068 W: AT BE WO 1989-US5067 US 1988-289,201 US 1989-315,736 A61K000-00 WO 1984-US2116 US 1983-566,172 G01N033-54 SYNTHETIQUES WO 8906138 WO 8502909 => d ab 36 LS ABEN LS AN TIEN TIFR PA LA DT PI DS AI PRAI PA LA DT PI DS AI PRAI LS AN TIEN TIFR CM Z

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Belain R; Wealner G J
Department of Internal Medicine, University of Iowa, Iowa City 52242, USA. lymphoma.

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Department of Immunology, University Hospital Utrecht, The Netherlands.
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Role of T-cell subsets in the **Dispecific antibody**Role of Rati-idiotype x anti-CD3) treatment of the BCL1 lymphoma.
Demanet C; Brissinck J; Lee O; Moser M; Thielemans K
Laboratory of Physiology, Medical School of the Vrije Universiteit Brussel Department of Internal Medicine, University of Iowa, Iowa City 52242, USA. LEUKEMIA AND LYMPHOMA, (1995 Jan) 16 (3-4) 199-207. Ref: 54 Journal code: 9007422. ISSN: 1042-8194. 9313244 PubMed ID: 768689
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de Gast G C; Haagen I A; van Houten A A; Klein S C; Duits A J; de Weger A; Vroom T M; Clark M R; Phillips J; van Dijk A J; +

Department of Immunology, University Hospital Utrecht, The Netherlands.

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General Review; (REVIEW)
(REVIEW, ACADEMIC)
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Role of T-cell subsets in the bispecific antibody
fanti-idiotype x anti-CD3) treatment of the BC11 lymphoma.
Demanet C; Brissinck J; Leo O; Moser M; Thielemans K
Laboratory of Physiology, Medical School of the Vrije Universiteit Brussell (VUB), Bellogum.
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Chuntharapai, Anan, 460 Point San Bruno Blvd., South San Francisco, CA, United States 94080 93315244 PubMed ID: 7686889
CD30-antigen-specific targeting and activation of T cells via murine bispecific monoclonal antibodies against CD3 and CD28: potential use for the treatment of Hodgkin's lymphoma.
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Pohl C: Denfeld R: Renner C; Jung W; Bohlen H; Sahin U; Hombach A; van Lier R; Schwonzen M; Diehl V; + Rohlen H; Sahin U; Hombach A; van Klinik I fur Innere Medizin, Universitat zu Koln, Cologne, Germany. INTERNATIONAL JOURNAL OF CANCER, (1993 Jul 9) 54 (5) 820-7. FUNCTIONAL STUDIES OF HUMAN IGG FC RECEPTORS. Institute for Cell and Developmental Biology Dartmouth Coll. Diss Abstr Int [B], (1991) 52 (3) 1335. ISSN: 0419-4217. United States Journal; Article; (JOURNAL ARTICLE) United States Journal; Article; (JOURNAL ARTICLE) Entered STN: 19950608 Last Updated on STN: 19950608 Entered STN: 19941107 Last Updated on STN: 19960517 ANSWER 13 OF 92 CANCERLIT 92679009 CANCERLIT CANCERLIT Entered STN: 19941107 Last Updated on STN: 19970509 CANCERLIT Entered STN: 19941107 Last Updated on STN: 19941107 English MEDLINE; Priority Journals MEDLINE 94243817 MEDLINE; Priority Journals MEDLINE 93315244 ANSWER 14 OF 92 USPATFULL ANSWER 12 OF 92 CAN ANSWER 11 OF 92 CA 94243817 CANCERLIT Erbe D V 92679009 (THESIS) English 199505 LL13 AN DN TI AU CS SO E NA DNA AU 20 SO SO CY DT LA FS OS EM ED DT LA EM EM ED ED

Hebert, Caroline, 460 Point San Bruno Blvd., South San Francisco, CA,

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Hybrid tryptophan aporepressor containing ligand binding sites
Lernhardt, Maldemar, Solana Basch, CA, United States
Bourdon, Mario, San Diego, CA, United States
Youderian, Phil, Ramona, CA, United States
Youderian, Phil, Ramona, CA, United States
(U.S. corporation)
19930302
US 5190873
US 1991-720222
19910621 (7)
Granted
Granted
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Carbohydrate-directed cross-linking reagents
IN Ashkenazi, Avi J., San Mateo, CA, United States
Chamow, Steven M., San Mateo, CA, United States
Kogan, Timothy P., Sugar Land, TX, United States
A Genentech, Inc., San Francisco, CA, United States
I US 5329028
I US 532907
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INCLS: 435/069.700, 435/069.100; 530/350.000; 530/812.000; 930/250.000
NCLM: 435/177.000
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ICM: CO7K016-28
ICS: CO7K016-24; C12N005-22
424/158.1; 530/388.73; 530/388.23; 530/389.2; 530/388.33; 530/389.1;
435/240.27
INDEXING IS AVAILABLE FOR THIS PATENT.
United States 94080
Kim, Kyung J., 460 Point San Bruno Blvd., South San Francisco, CA, United States 94080
Lee, James, 460 Point San Bruno Blvd., South San Francisco, CA, Un States 94080
US 5440021
US 5440021
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ICM: C07K013-00
ICS: C07K017-00; C07K017-02; C12P021-00
435/91; 435/69.7; 435/69.1; 435/177; 530/350; 530/812; 930/250
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INCLS: 530/388.230; 530/389.100; 530/389.200; 435/240.270
NCLM: 530/388.220
NCLS: 530/388.230; 530/389.100; 530/389.200
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548/546; 548/547; 548/548; 548/549
INDEXING IS AVAILABLE FOR THIS PATENT.
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PROCEDES SERVANT A PROVOQUER LA TOLERANCE DES LYMPHOCYTES T A UNE GREFEE
NOELLE, RANDOLPH, J.;
DURIE, Fiona, H.;
PARRER, David, C.;
APPEL, Michael, C.;
PHILIPS, Nancy, E.;
MORDES, JOHN, P. P.;
GRENIER, Dale, L.;
ROSSINI, Aldo, A.
TRUSTEES OF DARTMOUTH COLLEGE;
UNIVERSITY OF MASSACHUSETTS MEDICAL CENTER,
DURIE, FIONA, H.;
PARKER, David, C.;
PHILIPS, Nancy, E.;
MORDES, JOHN, P.;
PRIRER, David, C.;
PHILIPS, Nancy, E.;
MORDES, JOHN, P.;
GRENIER, Dale, L.;
PHILIPS, Nancy, E.;
MORDES, JOHN, P.;
GRENIER, Dale, L.;
ROSSINI, Aldo, A.
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199502846 PCTFULL ED 20020514
METHODS AND COPPOSITIONS FOR INHIBITING ENDOTHELIAL CELL AND FIBRINGGEN
METHODES ET COMPOSITIONS D'INHIBITION DE L'INFLAMMATION INDUITE PAR LES
ALTIERI, Dario, C.;
THORRING ENCYPELALES ET LE FIBRINGGNE
ALTIERI, DATIO, C.;
THORNION, GEOGGE, B.
THE SCRIPPS RESEARCH INSTITUTE
PAR 19951102
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BIOLOGICAL MARKERS OF BENIGN PROSTATE HYPERPLASIA MARQUEURS BIOLOGICALD BIL 'HYPERPLASIE BENIGNE DE LA IWRIGHT, George, L., Jr.
WRIGHT, George, L., Jr.
ENGLISH, George, L., Jr.
English
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ALDERSON, MASIE MAJEUR D'HISTOCOMPATIBILITE DE CLASSE II
ALDERSON, MASIE MAJEUR D'HISTOCOMPATIBILITE DE CLASSE II
ARMITAGE, Richard, J.;
COMEN, Jeffrey, I.;
COMEN, Jeffrey, I.;
FARRAH, Thereas, M.;
FARRAH, Thereas, M.;
FARRAH, Thereas, M.;
SPRIGGS, Melanie, K.;
IMMUNEX CORPORATION;
THE CURANORS OF THE UNIVERSITY OF MISSOURI;
NATIONAL INSTITUTES OF HEALTH
ENGLISH
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    WO 9533052
WO 9533052
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ISOLATED EPSTEIN-BARR VIRUS BZLF2 PROTEINS THAT BIND MHC CLASS II BETA
CHAINS
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AU CA JP AT BE CH DE DK ES FR GB GR IE IT LU 1
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IMMUNOSUPPRESSANT TARGET PROTEINS
BERLIN, Vivian;
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W: AU CA FI JP KR MX NO NZ AT BE 0
MC NL PT SE
WO 1995-US5348 A 19950428
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  FREEMAN, Gordon, J.;
NADLER, Lee, M.;
RENNERT, Paul;
JELLIS, Cindy, L.;
GREENFIELD, Edward;
GRAY, GATY, S.
REPLIGEN CORPORATION;
DANA FARBER CANCER INSTITUTE
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       CHIU, Marie, Isabel,
COTTAREL, Giullaume:
DAMAGNEZ, Veronique
MITOTIX, INC.
English
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US 1994-8/253,783
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1994-8/208,822 19940311 ANSWER 38 OF 92 PCTFULL COPYRIGHT 2003 Univentio
199502417 PCTFULL ED 20020514
NUCLECTIDE OR NUCLEOSIDE PHOTOAFFINITY COMPOUND MODIFIED
ANTIBODIES, METHODS FOR THEIR MANUFACTURE AND USE THEREOF
ANTICORES MODIFIES PAR DES COMPOSES DE PHOTOAFFINITE NUCLECTIDIQUES OU
HALEY, BOYA, E.;
KOHLER, Heinz;
RAJAGORALAN, Krishnan;
PAVLINKOVA, Gabriela
THE UNIVERSITY OF KENTUCKY RESEARCH FOUNDATION INHIBITORY AGENTS
PROCEDES AMELIORES DE TRANSPLANTATION A L'AIDE DE CELLULES MODIFIEES
D'AGENTS INHIBITEURS DE LYMPHOCYTES T
FRASER, Thomas
BIACRIN, INC.
English SE ANSWER 37 OF 92 PCTFULL COPYRIGHT 2003 Univentio 1995026740 PCTFULL ED 20020514 IMPROVED METHODS FOR TRANSPLANTATION USING MODIFIED CELLS AND T CELL SE SD SZ 1 W: --*,u42

MU CA JP AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT 1999-US4060

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MO 1995-US4060 ΡŢ 1995027042 PCTFULL COPYRIGHT 2003 Univentio 1995027042 PCTFULL ED 20020514 CEBRETICALLY MODIFIED CELLS FOR USE IN TRANSPLANTATION CELLULES GENETICALISMENT MODIFIEES EN VUE D'UNE TRANSPLANTATION ECAN E. Michael; CHAPPEL, Scott, C. DIACRIN, INC. Ä JP KE KG KP KR Z LK LR LT LU LV MD MG MN MW MX NG RO RU SD SE SG SI SK TJ TT UA UG US UZ VN KE MW SI BE CH DE DK ES PR GB GR IE IT LU MC NL PT SE BF BA WO 1995-US4024

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A 19950330

WI 1994-48/221,821

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COTKO16-18; C12N005-10; COTKO14-47; G01N033-577; A61K039-395; A61K038-13; A61K038-13; A61K035-39; A61K031-445; A61K035-39; A61K031-52; A61K035-39; A61K031-57; A61K035-39; WO 9526740 AU CA JP AT BE CH DE DK ES FR GB GR IE IT LU MC WW 1995-US3959 A 19950330 US 1994-8/220,724 19940331 US 1994-8/208,822 19940311 CO7K001-13 CO7K001-107; CO7K016-00; CO7K017-00; C12Q001-25; G01N033-53 A61K035-34; A61K048-00 Patent WO 9526740 Patent WO 9524417 W: A61K035-39 English L13 AN TIEN PA LA DT PI DS AI PRAI ICM AI PRAI ICM ICS LL13
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13	TE EE ES FI GB GE HU JP W MW NL NO NZ PL PT RO D SZ AT BE CH DE DE ES F CG CI CM GA GN ML MR	IAIRE (CEA) FR GB GR IE IT LU MC 2N015-13;	THE PROTEINS AND F POUR CES PROTEINES, EE ES FI GB GE HU JP EE ES FI GB GE HU JP SS ZAT BE CH DE DK CF CG CI CM GA GN ML
	48 OF 92 PCTFULL COPYRIGHT 2003 Univen 5976 PCTFULL ED 20020514 CRACIDS ENCODING A HOUSE DUST MITE ALLERGEN OR CALCISCES CODANT UN ALLERGENE D'ACARIENS NAYNE, R.; WAYNE, R.; KAW-YAN, A.; SICCE HARMACEUTICAL CORPORATION; UTE FOR CHILD HEALTH RESEARCH AM AT AU BB GG BB YC AC HC CZ DE D KE KG KP KR KZ LK LT LU LV MD MG M KE KG KP KR KZ LK LR LT LU LV MD MG M KE KG KP KR KZ LK LT LU LV MD MG M KE KG KP KR KZ LK LL LU LV MD MG M KR SO SE SI SK TJ TT UA UZ VN KE MW S FR GG RIE IT LU MC NL PT SE BF BJ C AUS14073 A 19991208	BER 49 OF 92 PCTFULL COPYRIGHT 2003 Univent	THER SO OF 92 PCTFULL COPYRIGHT 2003 Univent LISORA PCTFULL B. 20020514 L. APOPTOSIS-MODULATING PROTEINS, DNA ENCODING CODS OF USE THEREOF PROTEINS, DNA ENCODING CODS OF USE THEREOF PROTEINS MODULANT L'APOPTOSE, ADN CODAN EUR MACHAEL, C.; TER, MACHAEL, C.; "PHILID, J. "PHILIP, J. "PHILIP, J. "PHILIP, J. "PHILIP, J. "SER, MACHAEL, C.; "PHILIP, J. "PHILIP, J. "SER BE BY CA CH CN CZ DE DX NAT AU BB GG BR BY CA CH CN CZ DE DX KE CK FR EL LT LU W MD MG NN KE CK FR ER LT LT LU W MG NN FU SD SE SI SK TJ TT UA US UZ VN KE MR ES FR GB CR IE IT LU MC NL PT SE BF BJ MR NE SN TD TG

ANSWER 57 OF 92 PCTFULL COPYRIGHT 2003 Univentio
199401666 PCTFULL ED 20020513
ALLERGENIC PROTEINS AND PEPTIDES FROW DOG DANDER AND USES THEREFOR
PUTLISATEUR ASSOCIES
WORGENSTERN, Jay, P.;
KONIECZNY, Andrej;
BIZINKAUSKAS, Christine, B.;
BRADER, Andrew, W.
IMMULGOIC PHARMACEUTICAL CORPORATION;
WORGENSTERN, Jay, P.;
KONIECZNY, Andrej;
BIZINKAUSKAS, Christine, B.;
BRADER, Andrew, W.
RONIECZNY, Andrej;
BLISHAMSKAS, Christine, B.;
BRADER, Andrew, W.
ENDIROR ANDRESS, Christine, B.;
BRADER, Andrew, W.
BRADER, ANDRESS, Christine, B.;
BRADER, ANDRESS, Christine, B.; 1994015641 PCTFULL ED 20020513
METHOD FOR MODULATING TRANSENDOTHELIAL MIGRATION OF CELLS PROMOTING
INFLAMMATION, AND RELATED METHODS OF MEASUREMENT THEREOF
PROCEDE DE MODULATION DE LA MIGRATION TRANSENDOTHELIALE DES CELLULES
FAVORISANT L'INFLAMMATION ET PROCEDES CONNEXES DE MESURE DE CETTE 1994014976 PCTFULL COPYRIGHT 2003 Univentio 19940144976 PCTFULL ED 20020513 AND USES THEREOF IMMUNE RESPONSE MODULATIOR COMPLEX, AND USES THEREOF PIZZO, Salvatore, V.; CHU, Charleen, T.; OURY, Tim A61K037-02; G01N033-569; A61K039-395; A61K031-43; A61K039-395; A61K031-545; A61K039-395; A61K031-70 CI2N015-12 CI2P021-00; C07K015-06; C07K007-04; A61K039-35; C12P021-08 ES FR O LU MC NL E GB GR COPYRIGHT 2003 Univentio χÃ FR M: AL BB BG BR CA FI HU JP KP KR LK i AT BE CH DE DK ES FR GB GR IE IT I CI CM GA GN ML MR NE SN TD TG WO 1994-US416 A 19940112 AG1K039-395 19930112 BE CH DE I ES DK DK DE TJ TT UA US US UZ VN AT BE
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081,540
19930622 A2 19940721 KR US AT BE CH WO 1994-AU117 A 19940311 US 1993-8/031,141 19930312 US 1993-8/081,540 19930622 C12N015-12 GOTK013-00, AG1K039-35 A 19931230 19921231 19931122 MIGRATION MULLER, William, A. THE ROCKEFELLER UNIVERSITY; MULLER, William, A. PCTFULL OURY, Tim, D.
DUKE UNIVERSITY;
PIZZO, Salvatore, V.;
CHU, Charleen, T.; W: AU CA JP KE SE WO 1993-US12468 A US 1992-7/999,712 US 1993-8/156,549 ANSWER 58 OF 92 English Patent WO 9415641 WO 9416068 Patent AI PRAI ICM ICS L13 AN TIEN TIFR L13 AN TIEN L13 AN TIEN TIFR IN AI PRAI TIFR ICS ICM ICS NI ΡA N I I I IN LA DT PI DS PA ANSWER 56 OF 92 PCTFULL COPYRIGHT 2003 Univentio
394020614 PCTFULL ED 20020513
ALLERGENIC PROTEIN AND PEPTIDES FROM HOUSE DUST MITE AND USES THEREFOR
PROTEINE ET PEPTIDES ALLERGENES OBTENUS A PARTIR D'ACARIENS DETRITICOLES
ET LEURS UTILISATIONS
THOMAS, Wayne, Robert;
CHUA, KAW-Yan
INSTITUTE FOR CHILD HEALTH RESEARCH;
CHUA, KAW-Yan
CHUA, KAW-Yan
ENGlish ANSWER 5S OF 92 PCTFULL COPYRIGHT 2003 Univentio
1940.23 FO PCTFULL ED 20020513
TRANSGENIC ANIMAL MODEL FOR AUTOIMMUNE DISEASES
MODELE D'ANIMAL TRANSGENIQUE POUR MALADIES AUTOIMMUNES
MARLAN, DAVIG, M.;
JUNE, Carl, H.
THE UNITED STATES OF AMERICA as represented by THE SECRETARY OF THE NAVY XP SK JP KG 1 ES FR GB GR IE IT LU MC NL PT NL PT SE SE \mathbf{PT} Z, SD GB GE I ă Š US 1993-8/048.042 19930414
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AGIRO49-00; CO7H017-00; CI2N005-00; C12N015-00; G0IN031-00; E GB GR IE IT LU H FI PCTFULL COPYRIGHT 2003 Univentio Ξ ES GR DK ROSEN, Steven, D.;
SINGER, Mark, S.
GENBRYECH, INC.;
THE REGENCY OF CALIFORNIA;
LASKY, LAURENCE, A.;
BAUMHUETER, Susanne;
ROSEN, Steven, D.;
SINGER, Mark, S. A61K035-14; C12N005-20; C12P021-08; A61K037-02 A1 19940915 BB BG BR BY CA CH CN CZ DE I LK LU LV MD MG MN MW NL NO I В WO 9429436 A1 19941222 W: AU CA JP AT BE CH DE DK ES FR WO 1994-US6555 A 19940603 US 1993-8/073,223 19930604 WO 9423760 A1 19941027 W: AU CA JP AT BE CH DE DK ES FR W: 1994-US1674 A 19930414 US 1993-8/048,042 ANSWER 54 OF 92 PCTFULL COPYRIGHT 20
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INHIBITION OF LEUKOCYTE ADHESION DES LEUCOCYTES
INHIBITION DE L'ADHESION DES LEUCOCYTES MO 9425047 AU CA JP US AT BE CH DE DK IW 1994-US3791 A 19940406 US 1993-8/056,454 19930503 A61K039-395; C07K015-00 LASKY, Laurence, A ; BAUMHUETER, Susanne; REPLIGEN CORPORATION US 1993-8/073,223 C12N005-08 AU AT KR Patent WO 9423760 Patent WO 9420614 W: A61K037-02 English L13 AN TIEN TIFR IN L13 AN TIEN TIFR IN PA LA DT PI DS AI PRAI ICM L13 AN TIEN TIFR LA DT PI DS AI PRAI ICM LA DT PI DS AI PRAI ICM ΡA Z PA AT IA

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ΓA	OURY, Tim, D. English	AI PRAI	WO 1993-GB2492 A
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3	11-000000 (00000000000000000000000000000	TIFR	PROCEDE DE PREPARING SO
L13 AN	ANSWER 60 OF 92 PCTFULL COPYRIGHT 2003 Univentio 1994014467 PCTFULL ED 20020513	N	SPRIGGS, Melanie, K.; SRINIVASAN, Subhashini
TIEN	TREATMENT OF INFLAMMATORY BOWEL DISEASE WITH IFN-GAMMA INHIBITORS	PA	IMMUNEX CORPORATION
	INTERFERON GAMMA.	P. L.A	English Patent
N	ASHKENAZI, Avi, J.; WARD. Rebecca: H. R.	P.I.	9
PA	GENENTECH, INC.;	AI	W: AU CA JP N: WO 1993-US10034 A
	ASHKENAZI, Avi, J.; WARD, Rebecca, H., R.	PRAI	US 1992-7/969,703
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113	ANSWER 61 OF 92 PCTFULL, COPYRIGHT 2003 Univention		PICKER, Louis, J.;
AN	ULL ED 2002C	PA	BOARD OF REGENTS, THE
T FEN	MUDIIVALENI AND MUDIISFECIFIC BINDING PRUTEINS, THEIR MANUFACTURE AND USE	L'A	English Parent
TIFR	PROTEINES DE LIAISON MULTIVALENTES ET MULTISPECIFIQUES, LEUR FABRICATION ET LEUR HTILLGATION	PI	9409363
NI	Li BEON CITEDONILON HOLLIGER, Kaspar-Philipp;	DS AI	W: CA JP AT BI WO 1993-US9841 A
	GRIFFITHS, Andrew, David;	PRAI	072
	noosamoom, neiuricus, kenerus, Jacobus, marneus; MALMQVIST, Magnus;	ICS	G01N033-48 G01N033-533; G01N033-5;
	MARKS, James, David; McCHIMNES Drin Timothy.		
	POPE, Anthony, Richard;	L13 AN	ANSWER 64 OF 92 PCTF1
	PROSPERO, Terence, Derek;	TIEN	ANTIBODIES AGAINST TYPI
PA	MINIEK, GEGGOTY, BAUL CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED:	TIFR	ANTICORPS CONTRE LE REC
	MEDICAL RESEARCH COUNCIL;	:	GOEDDEL, David, V.;
	HOLLIGER, Kaspar-Philipp; GRIFFITHS, Andrew, David;		PALLADINO, Michael, A.
	HOOGENBOOM, Hendricus, Renerus, Jacobus, Matheus;	PA	
	MALMQVIST, Magnus; MARKS, James, David;	A F	English
	MCGUINNESS, Brian, Timothy;	PI	9137
	FUFE, Anthony, Kichard; PROSPERO, Terence, Derek:	DS	W: CA JP AT BI
Ą	WINTER, Gregory, Paul English	PRAI	US 1992-7/961,602
DI I		ICS	C12P021-08; C12N005-20
DS	AT AU BB BG BR BY CA CH CZ DE DK ES FI GB HU JP KP KR KZ		A61K035-14; G01N033-577
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THE SELECTION DIRECTE DE CELLULES PAR UN PRODUIT DE SECRETION MILTENT, SCEGAT, ANDERCAL, ANGERSE, ANDERCAL,

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OPPERMANN, Hermann CREATIVE BIOMOLECULES, INC.; CETUS ONCOLOGY CORPORATION CETUS ONCOLOGY CORPORATION CETUS ONCOLOGY CORPORATION CETUS ONCOLOGY CORPORATION CETUS CETUS A 19930819 W. 9316185 W. 9316185 A 19930205 US 1992-831,967 19920206 CIZNO15-62; CIZPOZI-08; CO7K015-28; CIZNO01-21; GOIN033-68; GOIN033-574	ANSWER 75 OF 92 1993011236 PCTFUI PRODUCTION OF ANT SEGMENT REPERTORING PRODUCTION D'ANTI REGIFFITHS, ANDREW HOOGENBOOM, HENDI MARKS, James, Daw MCAFFERTY, John;	GRIGG, GOOITEY, WALTER MEDICAL RESERVET COUNCIL: CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED; GRIFFITHS, Andrew, David; HOOGENBOOM, Hendricus, Renerus, Jacobus, Mattheus; MARKS, James, David; MCCAFFERTY, John; GRIGG, Geoffrey, Malter English	Parent WO 9311236 A1 W: AT AU BB BG MN MW NL NO GR IE IT LU TG WO 1992-GB2240 A		ASWERE 7 6 OF 92 PCTFULL COPYRIGHT 2003 Univentio 1993009803 PCTFULL ED 20020513 FACTOR X-DERIVED POLYPEPTIDES AND ANTI-PEPTIDE ANTIBODIES, SYSTEMS AND THERAREUTIC METHODS FOR THRIBITING INFLAMMATION POLYPEPTIDES DERIVES DU FACTEUR X ET ANTICORPS ANTIPEPTIDES, SYSTEMBS ET ALTIERI, Dario, C.; EDGINGTON, THOMAS, S.; ENCALPER SUSAn, C. +ef; THE SCRIPPS RESEBRCH INSTITUTE; THE SCRIPPS RESEBRCH INSTITUTE; EDGINTON, THOMAS, S. EDGINTON, THOMAS, S. EDGINTON, THOMAS, S. ENGLISH, DARIO, C.; EDGINTON, THOMAS, S. ENGLISH WO 9309803 AI 19930527
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COTK013-00; C12P021-00 ANSWER 78 OF 92 PCTFULL COPYRIGHT 2003 Univentio 1992006193 PCTFULL ED 20020513
ANTIBODIES DIRECTED AGAINST CD3
ANTICORES A EFFICACITE ANTICONISTE A L'ANTIGENE CD3
GORMAN, Scott, David;
WALDMANN, Herman
GORMAN, Scott, David;
WALDMANN, Herman
GORMAN, Scott, David;
WALDMANN, Herman
GORMAN, Herman
GORMAN, Scott, David;
EDG1ish 1993000356 PCTFULL COPYRIGHT 2003 Univentio 1993000356 PCTFULL ED 20020513 PTES PROTEINS CONTAINING BINDING SITES PROTEINES HYBRIDES RENFERMANT DES SITES DE LIAISON BOUNDON, MALGONA, VAUDERIAN PARTO, YOUDERIAN PARTO; ANSWER 79 OF 92 PCTFULL COPYRIGHT 2003 Univentio 1992006120 PCTFULL ED 20020513
ANTIGEN-ANTIBODY CONJUGATES
CONJUGUES ANTIGENES-ANTICORPS
MASON, Donald, William
MASON, Donald, William
English YOUDERIAN, Phil
CALIFORNIA INSTITUTE OF BIOLOGICAL RESEARCH;
LERNHARDT, Waldemar;
BOURDON, Mario;
YOUDERIAN, Phil MU 1991-061/20 A 19911004 GB 1990-9021679.7 19901005 C12N015-13 C07K015-28; C12P021-08; A61K039-395 A 19921120 19911122 WO 1992-US10068 US 1991-7/798,221 Patent WO 9206193 L13 AN TIEN TIFR IN L13 AN TIEN TIFR IN LA DT PI DS AI PRAI ICM LLA DT PI DS AI PRAI ICM L13 AN TIEN TIFR AI PRAI ICM ICS LA DT PI DS

ES W ÄΑ The Genuine Article (R) Number: TE709
PERFORMANCE OF CD3XCD19 BISPECIFIC MONOCLONAL-ANTIBODIES
IN B-CELL MALIGNANCY
HAAGEN I A (PEDIAL)
UNIV UTRECHT HOSP, DEPT IMMUNOL F03821, POSTBUS 85500, 3508 GA UTRECHT,
NETHERLANDS (Reprint) METHODS AND COMPOSITIONS FOR PROMOTING IMMUNOPOTENTIATION REACCEDES TO COMPOSITIONS DE PROMOTION DE L'IMMUNOPOTENTIALISATION BLUESTONE, JAFÉREY, A. ARCH DEVELOPMENT CORPORATION ÄΑ MG MG SE WO 9100360
W: AT AU BE CA CH DE DK ES FI FR GB IT JP LU NL NO :
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19890629
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C12N005-00; A61K039-395 W: AI 19910516
W: AT AT AU BB BE BF BG BJ BR CA CF CG CH CH CM DE ES FI FR GA GB GB CR HU IT JP KP KR LK LU LU MC NO SD SE SE SN SU TD TG
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PROHORMONE CLEAVAGE SITE BLOCKING ANTIBODY
ANTICORPS PROHORMONE POUR BLOCKING ANTIBODY
KRIEGLER, Michael;
PETEZ, Carl
CETUS CORPORATION COPYRIGHT 2003 Univentio COPYRIGHT 2003 Univentio WO 9102756 A1 18910307
W: AT AU BE CH DE DK ES FR GB IT JP LU NL WO 1990-US4536 A 19900813
US 1989-3955,254 19890816 BISPECTRIC REAGENTS FOR AIDS THERAPY
REACTIFS BISPECTFQUES POUR LE TRAITEMENT DU SIDA
FRANCER, Michael, W.;
GUYRE, Paul, M.;
DINNES, Nachan, B.
English ANSWER 86 OF 92 SCISEARCH COPYRIGHT 2003 ISI (R) 95:804167 SCISEARCH A61K039-395; G01N033-74; C12P021-08 PCTFULL ANSWER 85 OF 92 PCTFULL General Review; Journal LIFE; CLIN ANSWER 84 OF 92 Patent WO 9106319 W: Patent WO 9100360 NETHERLANDS English English ENGLISH AI PRAI L13 AN TIEN TIFR IN PA LA DT PI DS AI PRAI ICM L13 AN TIEN TIFR IN PA LA DT PI DS AI PRAI ICM ICM LL13 AN GA TI TI AU CS SO IN LA DT PI

No References Keyed

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

The Genuine Article (R) Number: RJ663
CDB T-CELLA CATIVATION AFTER INTRAVENCUS ADMINISTRATION OF CD3XCD19
BISPECIFIC ANTIBODA IN PATIENTS WITH
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NETHERLANDS; ENGLAND CANCER IMMUNOLOGY IMMUNOTHERAPY, (JUN 1995) Vol. 40, No. 6, pp. 390-396 The Genuine Article (R) Number: NN724
ROLE OF T-CELL SUBSETS IN TRE BISPECIFIC ANTIBODY
(ANTIDIOTYPE X ANTI-CL3) TREATHENT OF THE BCL(1) LYMPHOMA
DEMANET C; BRISSINCK J; LEE O; MOSER M; THIELEMANS K (REPTINC)
PREE UNT BRUSSELS, SCH MED, PHYSTOL LAB, LAARBEERLAAN 103-E, B-1090
BRUSSELS, BELGTUM (REPTINE); FREE UNIV BRUSSELS, SCH MED, PHYSTOL LAB, B-1090 BRUSSELS, BELGTUM; FREE UNIV BRUSSELS, B-1060 RHODE ST GENESE, WEINER G J (Reprint); DEGAST G C UNIV IOWA, DEPT INTERNAL MED, C32K GH, IOWA CITY, IA, 52242 (Reprint) LEUKEMIA & LYMPHOMA, (JAN 1995) Vol. 16, No. 3-4, pp. 199-207. ISSN: 1042-8194. General Review; Journal LIFE, CLIN CANCER RESEARCH, (01 JUN 1994) Vol. 54, No. 11, pp. 2973-2978. ISSN: 0008-5472. The Genuine Article (R) Number: MJ682
FYTOXINE RELEASE AFTER INTRAVENOUS ADMINISTRATION OF
BISPECIFIC ANTIBODY (CD3/CD19, SHR-1) - A PHASE-I STUDY
IN PATIENTS WITH CD19 POSITIVE B-CELL MALIGNANCIES Reference Count: 37 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* No References Keyed *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* Reference Count: 33 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* ANSWER 88 OF 92 SCISEARCH COPYRIGHT 2003 ISI (R) ANSWER 89 OF 92 SCISEARCH COPYRIGHT 2003 ISI (R) ANSWER 90 OF 92 SCISEARCH COPYRIGHT 2003 ISI (R) 93:729696 SCISEARCH SCISEARCH COPYRIGHT 2003 ISI (R) 95:97858 SCISEARCH The Genuine Article (R) Number: QC930 BISPECIFIC MONOCLONAL-ANTIBODY THERAPY OF B-ANSWER 87 OF 92 SCIS 95:495351 SCISEARCH SCISEARCH Article; Journal LIFE; CLIN ISSN: 0340-7004. Article; Journal CELL MALIGNANCY 94:329243 ENGLISH BELGIUM ENGLISH BELGIUM ENGLISH L13 AN GA TI CYA SO DT FS LA REC L13 GA DT FS LA REC L13 AN GA TI GAN 3 AU CS CYA SO DT FS LA REC CYA SO ΑU S AU

SEGAL D M (REPLIAL); QIAN J H; TITUS J A; MORENO M B; GEORGE A J T; JOST C R; KURUCZ I; ELGAMIL M; WUNDERLICH J R NIH, EXPTL IMMUNOL BRANCH, BLDG 10, ROOM 4B17, BETHESDA, MD, 20892 The Genuine Article (R) Number: LN208
The Genuine Article (R) Number: LN208
TO-SANTIGEN-SPECIFIC TARGETING AND ACTIVATION OF T-CELLS VIA MURINE
BISPECIFIC MONOCLONAL-ANTIBODIES AGAINST CD3 AND CD28
POTENTIAL USE FOR THE TREATHENT OF HODGKINS LYMPHONA
POLL C; DENFELD P; RENNER C; JUNG W; BOHLEN H; SHIN U; HOMBACH A; VANLIER
R; SCHWONZEN M; DIEHL V; PREUNDSCHUH M (REPTIN)
UNIV SAARLAND, MBD KILN I, W-6650 HOMBURG, GERMANY; UNIV COLOGNE, INNERE
MED KLIN I, W-5000 COLOGNE 41, GERMANY; NCB, AMSTERDAM, NETHERLANDS ADMINISTRATION OF BISPECIFIC ANTIBODY (CDJ/CDJ) 4. CDJ/CDJ 9. SHR.1) - A PHASE I STUDY IN PATIENTS WITH CD19 POSITIVE B-CELL MALIGNANCIES VANHOUTEN A R (REPTIAL); HAAGEN I A; CLARK M; GEERARS A; DELAU W; VROOM T M; BAST E J E G; DEGAST G C UNIV UTRECHT HOSP, DEPT HEMATOL, 3511 GV UTRECHT, NETHERLANDS; UNIV UTRECHT HOSP, DEPT IMMUNOL, 3511 GV VANHOUTEN A A (Reprint); HAAGEN I A; CLARK M; GEERARS A; DELAU W; VROOM M; BAST E J E G; DEGAST G C UNIVECTOR HOSP, DEPT HEMATOL, 3511 GV UTRECHT, NETHERLANDS; UNIV UTRECHT HOSP, DEPT HEMATOL, 3511 GV UTRECHT, NETHERLANDS; UNIV UTRECHT HOSP, DEPT PATHOL, 311 GV UTRECHT, NETHERLANDS; UNIV UTRECHT PATHOL, CAMBRIDGE, ENGLAND METHERLANDS; ENGLAND BLOOD, (15 NOV 1993) Vol. 82, No. 10, Supp. 1, pp. A580.

ISSN: 0006-4971. INTERNATIONAL JOURNAL OF CANCER, (09 JUL 1993) Vol. 54, No. 5, pp. INTERNATIONAL JOURNAL OF CANCER, (1992) Supp. 7, pp. 36-38. ISSN: 0020-7136. Article; Journal Reference Count: 32 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* Reference Count: 21 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* 93:729696 SCISEARCH MJ682 CYTOKINE RELEASE AFTER INTRAVENOUS ANSWER 92 OF 92 SCISEARCH COPYRIGHT 2003 ISI (R) COPYRIGHT 2003 ISI (R) SCISEARCH COPYRIGHT 2003 ISI (R) The Genuine Article (R) Number: JY980 TARGETED CYTOKINE PRODUCTION SCISEARCH => d 90, 88, 86, 82, 61 ibib ab GERMANY; NETHERLANDS SCISEARCH SCISEARCH Conference; Journal LIFE; CLIN ISSN: 0020-7136. Article; Journal L13 ANSWER 90 OF 92 ANSWER 91 OF 92 THE GENUINE ARTICLE: No References ACCESSION NUMBER: CORPORATE SOURCE: 92:687393 (Reprint) ENGLISH ENGLISH 820-827 ENGLISH L13 AN GA DT FS LA REC CYA DT FS LA REC LL13 AN GA TI CYA SO DT FS LA REC ΑN S SS S

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**REFERENCE COURT: **ABSTRACT IS A LUMOR ASSOCIATED WITH the other arm can retarget T-cells toward tumor cells in an MHC independent manner, the power of the cellular immune system. B-cell an MHC independent manner, the power of the cellular immune system. B-cell antibody therapy because of their sensitivity to other forms of antibody therapy, and the extent to which B-cells and T-cells communicate at the molecular level. BsAbs that recognize CD3 and a number of antigens on malignant B-cells and models of antigens on malignant B-cells in animal models of B-cell malignancy back and an undersome of antigens on malignant to unmodified monoclonal antibody therapy. Organis start to be capable of retargeting T-cells. In animal models of B-cell malignancy back therapy of significant biologic effects, and suggest they have anti-tumor activity as well. A number of significant duestions relating to babb therapy of endogenously produced and exogenously administered cytokines are likely to play. Further exploration of whether bsab can induce T-cells to target to tumor will also be required before the true promise of this proved.
                            DEPT PATHOL, 3511
DEPT PATHOL,
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OF B-CELL MALIGNANCY
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UNIV IOWA, DEPT INTERNAL MED, C32K GH, IOWA CITY,
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GV UTRECHT, NETHERLANDS; UNIV UTRECHT HOSP, GV UTRECHT, NETHERLANDS; UNIV CAMBRIDGE, 1 CAMBRIDGE, ENGLAND NETHERLANDS; ENGLAND BLOOD, (15 NOV 1993) Vol. 82, No. 10, Supp EISN: 0006-4971.
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THE GENUINE ARTICLE: QC930
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*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS.

*Bispecific monoclonal antibodies, with a dual specific monoclonal antibodies, with a dual arteros on immune effector cells, have been shown (in vitro) to be effective in directing and triggering effector cells to kill target cells resulting in target cell lysis. Bispecific monoclonal antibodies (BsAb) against the CD3 antigen on T cells and the CD19 antigen on B call were developed. Data obtained by in vitro experiments might indicate that clinical responses in BsAb immunotherapy, will only be obtained in patients with minmal tumor load and may need additional T cell sminlation via cytofkines such as IL-2. Although these experiments have shown us their limitations, they also include the promise of BsAb-directed immunotherapy in B coll mailgnancy as further demonstrated during a plasse I trail, showing little toxicity. Clearly, much remains to be done before this showing little toxicity. Clearly, much remains to be done before this showing little toxicity. Clearly, much remains to be done before this coll malignancy. This report describes the experiments performed to test a new immunotherapeutic agent in B cell malignancy. Bisspecific antibodies are described that can target cycloxic T cells to tumor cells and elicit a cytolytic action towards these cancer cells. ENGLISH REFERENCE COUNT: FILE SEGMENT: LANGUAGE:

2 PCTFULL COPYRIGHT 2003 Univentio 1991008774 PCTFULL ED 20020513 CTTOKINE ANTIBODY FOR THE TREATMENT OF SEPSIS ANTICORRS DE CYTOKINE UTILISE DANS LE TRAITEMENT DE LA SEPTICEMIE ARDEN, Lucien, A.; CREASEY, ADIA, A.; KRASEY, ALIA, A.; KRASEY, KISTON, E. KOTHS, KISTON, E. CETUS CORPORATION; AARDEN, Lucien, A. English Patent L13 ANSWER 82 OF 92 ACCESSION NUMBER: PATENT ASSIGNEE(S): LANGUAGE OF PUBL.: DOCUMENT TYPE: PATENT INFORMATION: TITLE (ENGLISH): TITLE (FRENCH): INVENTOR(S):

Al 19910627 WO 9108774 DESIGNATED STATES

KIND

NUMBER

APPLICATION INFO.

WE APPLICATION INFO.

WO 1990-US7411

A 19901213

PRIORITY INFO.:

US 1989-451,218

19891215

ABEN

Compositions and methods for prophlactically or therapeutically

treating sepsis consisting of

antibody to IL-6 and/or M-CSF wherein the antibodies

are administered alone or in combination.

ABER

Compositions et procedes de traitements prophylatiques ou therapeutiques

comprenant un anticorps contre IL-6 et/ou les facteurs de croissance de macrophages, dans lesquels $% \left(\frac{1}{2}\right) =\frac{1}{2}\left(\frac{1}{2}\right) +\frac{1}{2}\left(\frac{1}{2}\right$ de la septicemie,

les anticorps sont administres seuls ou de maniere combinee.

1994013804 PCTFULL ED 20020513 MULTIVALENT AND MULTISPECIFIC BINDING PROTEINS, THEIR MANUFACTURE AND USE COPYRIGHT 2003 Univentio ANSWER 61 OF 92 ACCESSION NUMBER: TITLE (ENGLISH):

PROTEINES DE LIAISON MULTIVALENTES ET MULTISPECIFIQUES, LEUR FABRICATION ET LEUR UTILISATION HOLLIGABAL-PHILIDP; GRIFFITHS, Andrew, David; INVENTOR(S):

TITLE (FRENCH):

Polypeptides comprising a first domain, which comprises a binding region RA E JP KP SE SK U HOLLIGER, Kaspar-Philipp;

GRIPFITHS, Andrew, David;

GROGENBOOM, Hendricus, Renerus, Jacobus, Matheus;

MALMOVIST, Magnus;

MARKS, James, David;

MARKS, Annes, David;

POPE, Anthony, Richard;

POPE, Anthony, Richard;

POPE, Anthony, Richard;

MINTER, Gregory, Paul HOOGENBOOM, Hendricus, Renerus, Jacobus, Matheus; 35835 88 FT CT FI RO IE SN MCGUINNESS, Brian, Timothy;
POPE, Anthony, Richard;
POSOSERO, Terence, Derek;
WINTER, Gregory, Paul
CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED;
MEDICAL RESEARCH COUNCIL; AT AU BB BC BR BY CA CH CZ DE DK ES FI KZ LL UL UW MG MN MN LN ON Z PL PT RO US UZ VN AT BE CH DE DK ES FR GB RI E SE BF BJ CF CG CI CM GA GN ML MR NE SN WO 1999-GB2492 A 199311203 GB 1993-9325453.1 19931014 GB 1993-9310816.7 19930510 GB 1993-9310816.7 19930512 GB 1993-9310816.7 19930512 DATE Al 19940623 KIND David; MALMQVIST, Magnus Јашев, WO 9413804 WINTER, English NUMBER Patent PATENT ASSIGNEE(S): LANGUAGE OF PUBL.: DOCUMENT TYPE: PATENT INFORMATION: APPLICATION INFO.: PRIORITY INFO.: DESIGNATED STATES ABEN

of an immunoglobulin heavy chain variable region, and a second domain, which comprises a binding region of an immunoglobulin light chain variable region, the domains being linked but immunoglobulin light chain variable region, the domains being linked but with each other to form an antigen binding site, associate to form with each other to form any be multiwalent or have multispecificity. The domains may be linked by a short peptide linker or may be joined directly together. Bispecific dimers may have longer linkers.

whethods of preparation of the polypeptides and multimers and diverse repertoires thereof, and their display on the surface of bacteriophage for easy selection of binders of

interest, are disclosed, along with many utilities.

Des polypeptides comprenant un premier domaine presentant une region de fixation de la region variable d'une chaine lourde d'immunoglobuline, et un deuxieme domaine d'antigene, s'associent pour former des multimeres de fixation d'antigene tels que des dimeres, lesquels peuvent etre multivalents ou a specificite multiple. Les domaines peuvent etre lies par un lieur peptidique court ou unis directement. Les dimeres bispecifiques peuvent avoir des lieurs plus directement. Les dimeres bispecifiques peuvent avoir des lieurs plus fixation de la region variable d'une chaine legere d'immunoglobuline, lesdits domaines etant lies mais incapables de s'associer pour former un site de fixation presentant une region de Des procedes de ABFR

preparation desdits polypeptides et multimeres ainsi que de leurs divers regertoires, leur utilisation a la surface d'un bacteriophage afin de faciliter la selection des lieurs selon l'invention ainsi que leurs, nombreuses applications sont decrits.

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Connection closed by remote host

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